

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:27:20 ; Search time 8.32 Seconds  
(without alignments)  
150.299 Million cell updates/sec

Title: US-09-786-214A-14

Perfect score: 65

Sequence: 1 LPVVGLSPGEQE 13

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 78:\*  
1: Pirl1.\*  
2: Pirl2.\*  
3: Pirl3.\*  
4: Pirl4.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	44	67.7	381	2	F75270 cytochrome P450 -
2	43	66.2	821	2	C84304 DNA helicase [impo
3	41	63.1	156	2	F87551 conserved hypothet
4	41	63.1	326	2	T45226 probable N5,N10-me
5	40	61.5	243	2	I54459 MHC H-2K1-k - mous
6	40	61.5	428	2	AG1304 uracil permease ho
7	40	61.5	428	2	AG1676 uracil permease ho
8	40	61.5	540	2	A75250 carboxylesterase,
9	40	61.5	623	2	T40991 probable lysophosp
10	39	60.0	43	2	S21065 Ig kappa chain V r
11	39	60.0	96	2	S45441 Ig kappa chain V r
12	39	60.0	103	2	S19975 Ig kappa chain V r
13	39	60.0	106	2	P80070 Ig kappa chain V r
14	39	60.0	106	2	P4282 Ig kappa chain (an
15	39	60.0	107	2	S57444 Ig kappa chain V-J
16	39	60.0	108	2	C30502 Ig kappa chain V r
17	39	60.0	108	2	S33988 Ig kappa chain V r
18	39	60.0	108	2	G44151 Ig kappa chain V r
19	39	60.0	111	2	S23628 Ig kappa chain V r
20	39	60.0	114	2	S34905 Ig kappa chain V r
21	39	60.0	115	1	K3HUVG Ig kappa chain pre
22	39	60.0	115	1	KVMSL7 Ig kappa chain pre
23	39	60.0	115	2	S11697 Ig kappa chain pre
24	39	60.0	116	2	B25521 Ig kappa chain pre
25	39	60.0	119	2	S41816 Ig kappa chain V r
26	39	60.0	125	2	S40344 Ig kappa chain V-J
27	39	60.0	128	2	PNO445 Ig kappa chain pre
28	39	60.0	128	2	S40379 Ig kappa chain V-J
29	39	60.0	128	2	A56701 Ig kappa chain V r

30 39 60.0 129 2 S29627 Ig kappa chain V r  
31 39 60.0 129 2 S40363 Ig kappa chain - h  
32 39 60.0 132 2 S05268 Ig kappa chain pre  
33 39 60.0 144 2 PLO106 Ig kappa chain pre  
34 39 60.0 144 2 B30502 Ig heavy chain V r  
35 39 60.0 215 2 A23746 Ig kappa chain V-I  
36 39 60.0 319 2 AD0941 probable regulator  
37 39 60.0 508 2 E70764 probable cobi prot  
38 39 60.0 563 2 JQ0623 nerve growth facto  
39 39 60.0 1367 2 S74285 BUD3 protein - yea  
40 39 60.0 1506 2 JCS985 phosphoinositide 3  
41 38.5 59.2 656 2 E75468 hypothetical prote  
42 38 58.5 86 2 S20649 Ig heavy chain V r  
43 38 58.5 110 2 S60591 Ig light chain var  
44 38 58.5 111 2 PNO537 Ig kappa chain V r  
45 38 58.5 152 2 G75184 probable transcrip

#### ALIGNMENTS

##### RESULT 1

F75270  
cytochrome P450 - Deinococcus radiodurans (strain R1)  
C:Species: Deinococcus radiodurans  
C:Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000  
C:Accession: F75270  
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;  
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; M.  
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.  
Science 286, 1571-1577, 1999  
A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.  
A:Reference number: A75250; MUID:20036896; PMID:10567266  
A:Accession: F75270  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-381 <WHI>  
A:Cross-references: GB:AE002076; GB:AE000513; NID:96460285; PIDN:AAF12016.1; PID:964602  
A:Experimental source: strain R1  
C:Genetics:  
A:Gene: DR2473  
A:Map position: 1

Query Match 67.7% Score 44; DB 2; Length 381;  
Best Local Similarity 100.0%; Pred. No. 8.7; Indels 0; Gaps 0;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 LPVVVGLSP 9

Db 52 LPVVVGLSP 60

##### RESULT 2

C84304  
DNA helicase [imported] - Halobacterium sp. NRC-1  
C:Species: Halobacterium sp. NRC-1  
C:Date: 02-Feb-2001 #sequence\_revision 02-Feb-2001 #text\_change 02-Feb-2001  
C:Accession: C84304  
R:Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, J.  
Leithauser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jabl  
Jung, K.H.; Alam, M.; Freitas, T.  
Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000  
A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; L.  
A:Title: Genome sequence of Halobacterium species NRC-1.  
A:Reference number: A84160; MUID:20504483; PMID:11016950  
A:Accession: C84304  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-821 <STO>  
A:Cross-references: GB:AE004437; NID:gl0580995; PIDN:AAG19799.1; GSPDB:GN00138  
C:Genetics:  
A:Gene: hel

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Query Match      66.2%; Score 43; DB 2; Length 821;
Best Local Similarity 90.0%; Pred. No. 28;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGQEQ 12
   |||||
Db 326 AVVGLSPAEQ 335

RESULT 3
F87551
conserved hypothetical protein CC2439 [imported] - Caulobacter crescentus
C:Species: Caulobacter crescentus
C:Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 10-May-2001
C:Accession: F87551
R:Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolon
n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M.
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A:Title: Complete Genome Sequence of Caulobacter crescentus.
A:Reference number: AB7249; MUID:21173698; PMID:11259647
A:Accession: F87551
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-156 <STO>
A:Cross-references: GB:AE005673; NID:gl3423984; PIDN:AAK24410.1; GSPDB:GN00148
C:Genetics:
C:Superfamily: Haemophilus influenzae conserved hypothetical protein HI0305

Query Match      63.1%; Score 41; DB 2; Length 156;
Best Local Similarity 80.0%; Pred. No. 12;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGQEQ 12
   |||||
Db 18 AVVGLDPGEK 27

RESULT 4
T45226
probable NS10-methylene-tetrahydrodethanopterin reductase (F420-dependent) [imported]
C:Species: Methanobolus tindarius
C:Date: 21-Jan-2000 #sequence_revision 21-Jan-2000 #text_change 21-Jan-2000
C:Accession: T45226
R:Westenberg, D.J.; Braune, A.; Ruppert, C.; Mueller, V.; Herzberg, C.; Gottschalk, G.;
submitted to the EMBL Data Library, September 1998
A:Description: The F420H2-dehydrogenase from Methanobolus tindarius: Cloning of the ffd
A:Reference number: Z22947
A:Accession: T45226
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-326 <WES>
A:Cross-references: EMBL:AJ011519; PIDN:CAB56639.1
A:Experimental source: DSM 2278
C:Genetics:
A:Gene: ffdA

Query Match      63.1%; Score 41; DB 2; Length 326;
Best Local Similarity 70.0%; Pred. No. 24;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGQEQ 12
   |||||
Db 88 AILGLPGGEQ 97

RESULT 5
I54459
MHC H-2K1-k - mouse
C:Species: Mus musculus (house mouse)
C:Date: 02-Aug-1996 #sequence_revision 02-Aug-1996 #text_change 05-Nov-1999
C:Accession: I54459

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R:Watts, S.; Davis, A.C.; Goodenow, R.S.
Immunogenetics 29, 355-357, 1989
A:Title: Sequence analysis of the C3H H-2K1-k gene: Relationship to the H-2 loci.
A:Reference number: I54459; MUID:89233303; PMID:2714856
A:Accession: I54459
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-243 <RES>
A:Cross-references: GB:M27134; NID:gl199435; PIDN:AAA39610.1; PID:g387456
C:Genetics:
A:Introns: 22/1; 112/1

Query Match      61.5%; Score 40; DB 2; Length 243;
Best Local Similarity 77.8%; Pred. No. 27;
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 5 VGLSPGQEQ 13
   |||||
Db 216 LGLSPGEE 224

RESULT 6
AG1304
uracil permease homolog pyrP [imported] - Listeria monocytogenes (strain EGD-e)
C:Species: Listeria monocytogenes
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AG1304
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloecke
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, H
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kref, J.; Kuhn, M.; Kunst, F.; Kurapat, G.; Madueno, E.; Maitournam, A.; M
ok, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wetland
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669
A:Accession: AG1304
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-428 <GLA>
A:Cross-references: GB:NC_003210; PIDN:CAC99917.1; PID:gl16411293; GSPDB:GN00177
A:Experimental source: strain EGD-e
C:Genetics:
A:Gene: pyrP
C:Superfamily: uracil transport protein uraA

Query Match      61.5%; Score 40; DB 2; Length 428;
Best Local Similarity 70.0%; Pred. No. 47;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 LPAYVGLSPG 10
   |||||
Db 42 VPSVTGLSPG 51

RESULT 7
AG1676
uracil permease homolog pyrP [imported] - Listeria innocua (strain Clip11262)
C:Species: Listeria innocua
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AG1676
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloecke
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, H
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kref, J.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wetland
ok, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wetland
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669
A:Accession: AG1676
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-428 <GLA>
A:Cross-references: GB:AL592022; PIDN:CAC97183.1; PID:gl16414454; GSPDB:GN00178
A:Experimental source: strain EGD-e

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A;Experimental source: strain Clip11262

C;Genetics:

A;Gene: pyrP

C;Superfamily: uracil transport protein uraA

Query Match 61.5%; Score 40; DB 2; Length 428;  
Best Local Similarity 70.0%; Pred. No. 47;  
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 LPAVVGSLSPG 10

Db 42 VPSVTGLSPG 51

RESULT 8

A75250

carboxylesterase, type B - Deinococcus radiodurans (strain R1)

C;Species: Deinococcus radiodurans

C;Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000

C;Accession: A75250

R;White, O.; Eissen, J.A.; Heideberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;

M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; Ma

S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.

Science 286, 1571-1577, 1999

A;Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.

A;Reference number: A75250; MUID:20036896; PMID:10567266

A;Accession: A75250

A;Molecule type: DNA

A;Residues: 1-540 <WHI>

A;Cross-references: GB:AE002092; GB:AE000513; NID:g6460455; PIDN:AAF12163.1; PID:g646045

A;Experimental source: strain R1

C;Genetics:

A;Gene: DR2626

A;Map position: 1

C;Superfamily: cholinesterase; cholinesterase homology

Query Match 61.5%; Score 40; DB 2; Length 540;  
Best Local Similarity 70.0%; Pred. No. 60;  
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 PAVVGLSPGE 11

Db 512 PQVGLTAPGE 521

RESULT 9

T40991

probable lysophospholipase precursor - fission yeast (Schizosaccharomyces pombe)

C;Species: Schizosaccharomyces pombe

C;Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 21-Jan-2000

C;Accession: T40991

R;Lyne, M.; Rajandream, M.A.; Barrell, B.G.; Volckaert, G.

submitted to the EMBL Data Library, March 1999

A;Reference number: Z21962

A;Accession: T40991

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-623 <LYN>

A;Cross-references: EMBL:AL049559; PIDN:CNB40176.1; GSPDB:GN000668; SPDB:SPCC1450.09C

A;Experimental source: strain 972h; cosmid c1450

C;Genetics:

A;Gene: SPDB:SPCC1450.09C

A;Map position: 3

C;Superfamily: yeast lysophospholipase

Query Match 61.5%; Score 40; DB 2; Length 623;  
Best Local Similarity 75.0%; Pred. No. 69;  
Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 PAVVGLSPGE 13

Db 76 PASDGLSTGE 87

RESULT 10

S21065

Ig kappa chain V region (anti-RH(D)) - human (fragment)

C;Species: Homo sapiens (man)

C;Date: 22-Nov-1993 #sequence\_revision 24-May-1996 #text\_change 09-May-1997

C;Accession: S21065

R;Plouha, A.; Lecroisey, A.; Henschen, A.; Rouger, P.; Keil, B.

Protein Seq. Data Anal. 4, 317-318, 1991

A;Title: Subgroup assignment of a human monoclonal anti-Rh(D) antibody.

A;Reference number: S21065; MUID:92253544; PMID:1812483

A;Accession: S21065

A;Molecule type: protein

A;Residues: 1-43 <DLO>

C;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

Query Match 60.0%; Score 39; DB 2; Length 43;  
Best Local Similarity 63.6%; Pred. No. 69;  
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGE 12

Db 8 PATLSLSPGER 18

RESULT 11

S45441

Ig kappa chain V region - human (fragment)

C;Species: Homo sapiens (man)

C;Date: 27-Jan-1995 #sequence\_revision 27-Jan-1995 #text\_change 21-Jan-2000

C;Accession: S45441

R;Cox, J.P.L.; Tomlinson, I.M.; Winter, G.

Eur. J. Immunol. 24, 827-836, 1994

A;Title: A directory of human germ-line V(kappa) segments reveals a strong bias in their

A;Reference number: S45324; MUID:94200218; PMID:8149953

A;Accession: S45441

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-96 <COX>

A;Cross-references: EMBL:Z27500

C;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

F;16-91/Domain: immunoglobulin homology <IMM>

Query Match 60.0%; Score 39; DB 2; Length 96;  
Best Local Similarity 63.6%; Pred. No. 15;  
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGE 12

Db 8 PATLSLSPGER 18

RESULT 12

S19975

Ig kappa chain V region (M-T408) - mouse (fragment)

C;Species: Mus musculus (house mouse)

C;Date: 06-Feb-1995 #sequence\_revision 06-Feb-1995 #text\_change 21-Jan-2000

C;Accession: S19975

R;Weissenhorn, W.; Riethmuller, G.; Weiss, E.M.; Rieber, E.P.

submitted to the EMBL Data Library, March 1992

A;Description: Structural characterization of CD4 mAb.

A;Reference number: S19963

A;Accession: S19975

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-103 <WFI>

A;Cross-references: EMBL:X65097; NID:g52296; PIDN:CAA46225.1; PID:g52297

C;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

F;11-85/Domain: immunoglobulin homology <IMM>

Query Match 60.0%; Score 39; DB 2; Length 103;  
 Best Local Similarity 63.6%; Pred. No. 17;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
 QY 2 PAVVGLSPGEQ 12  
 ||:|||||:  
 Db 3 PATLSLSPGER 13

## RESULT 13

Ig kappa chain V region (38C13.V6.1) - mouse  
 C:Species: Mus musculus (house mouse)  
 C>Date: 31-Mar-1990 #sequence\_revision 31-Mar-1990 #text\_change 21-Jan-2000  
 C:Accession: PS0070  
 R;Levy, S.; Campbell, M.J.; Levy, R.  
 J. Exp. Med. 170, 1-13, 1989  
 A;Title: Functional immunoglobulin light chain genes are replaced by ongoing rearrangement  
 A;Reference number: A92781; MUID:89310348; PMID:2501443  
 A;Accession: PS0070  
 A;Status: translation not shown  
 A;Molecule type: DNA  
 A;Residues: 1-106 <LEV>  
 C;Superfamily: immunoglobulin V region; immunoglobulin homology  
 C;Keywords: heterotetramer; immunoglobulin  
 F;16-89/Domain: immunoglobulin homology <IMM>

Query Match 60.0%; Score 39; DB 2; Length 106;  
 Best Local Similarity 54.5%; Pred. No. 17;  
 Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 ||:|||||:  
 Db 8 PATIAAGSGEK 18

## RESULT 14

PC4282  
 Ig kappa chain (anti-ss-A/Ro 60K peptide) (E-42 and E-56) - human (fragment)  
 C:Species: Homo sapiens (man)  
 C>Date: 07-Jul-1997 #sequence\_revision 29-Aug-1997 #text\_change 21-Jan-2000  
 C:Accession: PC4282; PC4284  
 R;Suzuki, H.; Takemura, H.; Suzuki, M.; Sekine, Y.; Kashiwagi, H.  
 Biochem. Biophys. Res. Commun. 232, 101-106, 1997  
 A;Title: Molecular cloning of anti-ss-A/Ro 60-kDa peptide fab fragments from infiltrating  
 A;Reference number: PC4279; MUID:97236289; PMID:9125110  
 A;Accession: PC4282  
 A;Molecule type: protein  
 A;Residues: 1-106 <SU2>  
 A;Note: E-42  
 A;Accession: PC4284  
 A;Molecule type: protein  
 A;Residues: 1-106 <SU2>  
 A;Note: E-56  
 C;Comment: This antibody is commonly found in systemic autoimmune diseases such as Sjogren  
 C;Superfamily: immunoglobulin V region; immunoglobulin homology  
 F;14-88/Domain: immunoglobulin homology <IMM>

Query Match 60.0%; Score 39; DB 2; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 17;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 ||:|||||:  
 Db 6 PATLSLSPGER 16

## RESULT 15

SS7444  
 Ig kappa chain V-J region - human (fragment)  
 C:Species: Homo sapiens (man)  
 C>Date: 10-Oct-1995 #sequence\_revision 17-Nov-1995 #text\_change 21-Jan-2000  
 C:Accession: SS7444

R;Paterson, G.; Wilson, G.; Kennedy, P.G.E.; Willison, H.J.

submitted to the EMBL Data Library, June 1995

A;Description: Analysis of anti-GM1 ganglioside IgM antibodies cloned from motor neurop

A;Reference number: SS7408

A;Accession: SS7444

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-107 <PAT>

A;Cross-references: EMBL:X87898; NID:g871275; PIDN:CAA61149.1; PID:g871276

C;Superfamily: immunoglobulin V region; immunoglobulin homology

F;16-90/Domain: immunoglobulin homology <IMM>

Query Match 60.0%; Score 39; DB 2; Length 107;  
 Best Local Similarity 63.6%; Pred. No. 17;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 ||:|||||:  
 Db 8 PATLSLSPGER 18

Search completed: May 7, 2004, 12:39:07  
 Job time : 8.48667 secs



GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 4.68 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-13

Perfect score: 68

Sequence: 1 PAVVGLSPGEQY 13

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	43	63.2	211	1	COBL_METJUA
2	43	63.2	633	1	PLB5_SCHPO
3	41	60.3	156	1	RUVX_CAUCR
4	41	60.3	326	1	MER_METTI
5	41	60.3	508	1	COBI_MYCTU
6	40	58.8	673	1	PLB4_SCHPO
7	39	57.4	115	1	KV31_HUMAN
8	39	57.4	115	1	KV51_MOUSE
9	39	57.4	1636	1	BUD3_YEAST
10	38	55.9	429	1	RS1_LEULA
11	38	55.9	507	1	CAT3_PICAN
12	37	54.4	446	1	COB3_ARCFU
13	37	54.4	446	1	ENOL1_MALZE
14	37	54.4	597	1	NR41_RAT
15	37	54.4	607	1	GLMS_CLOTE
16	37	54.4	813	1	CAD3_MOUSE
17	37	54.4	1402	1	NI60_MOUSE
18	36.5	53.7	1121	1	BMS1_SCHPO
19	36	52.9	129	1	KV3H_HUMAN
20	36	52.9	158	1	RSD_ECOLI
21	36	52.9	162	1	RSD_SALTY
22	36	52.9	243	1	SMT1_SYNP7
23	36	52.9	279	1	TSY3_CAUCR
24	36	52.9	317	1	OADI_HUMAN
25	36	52.9	342	1	HUPK_AZOVI
26	36	52.9	348	1	HOKV_AZOVI
27	36	52.9	422	1	TKSU_PYRKO
28	36	52.9	464	1	LEU2_BACCR
29	36	52.9	471	1	CD36_BOVIN
30	36	52.9	590	1	MUTL1_THEIN
31	36	52.9	609	1	GLMS_CLOPE
32	36	52.9	614	1	CPRI_DROME
33	36	52.9	624	1	PLB2_SCHPO

34	36	52.9	699	1	EFG_HAFIN
35	36	52.9	700	1	EFG_PASMU
36	36	52.9	750	1	ELS_CHICK
37	36	52.9	844	1	HEXA_STRPN
38	36	52.9	926	1	PTM4_HUMAN
39	36	52.9	1103	1	VG37_BPARI
40	36	52.9	1173	1	ATC2_YEAST
41	36	52.9	3703	1	ABF1_HUMAN
42	35	51.5	100	1	KV3C_HUMAN
43	35	51.5	109	1	KV3B_HUMAN
44	35	51.5	109	1	KV3D_HUMAN
45	35	51.5	109	1	KV3E_HUMAN

## ALIGNMENTS

### RESULT 1

ID	COBL_METJUA	STANDARD;	PRT;	211 AA.
AC	Q58917;			
DT	15-DEC-1998 (Rel. 37, Created)			
DT	15-DEC-1998 (Rel. 37, Last sequence update)			
DT	28-FEB-2003 (Rel. 41, Last annotation update)			
DE	Probable precorrin-6Y C5,15-methyltransferase [decarboxylating] (BC 2.1.1.132) (Precorrin-6 methyltransferase) (Precorrin-6Y methylase).			
DE	COBL OR MJ1522.			
GN	Methanococcus jannaschii.			
OS	Archaea; Euryarchaeota; Methanococci; Methanococcales;			
OC	Methanocaldococcaceae; Methanocaldococcus.			
OX	NCBI_TaxID=2190;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN=JAL-1 / DSM 2661 / ATCC 43067;			
RX	MEDLINE=96337999; PubMed=8689087;			
RA	Bult C.J., White O., Olsen G.J., Zhou L., Fleischmann R.D., Sutton G.G., Blake J.A., FitzGerald L.M., Clayton R.A., Gocayne J.D., Kerlavage A.R., Dougherty B.A., Tomb J.-F., Adams M.D., Reich C.I., Overbeek R., Kirkness E.F., Weinstock K.G., Merrick J.M., Glodek A., Scott J.B., Geoghegan N.S.M., Weidman J.F., Sadow P.W., Hanna M.C., Uutterback T.R., Kelley J.M., Peterson J.D., Klenk H.-P., Roberts K.M., Hurst M.A., Kaine B.P., Borodovsky M., Klenk H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;			
RT	*Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii*.			
RL	Science 273:1058-1073 (1996).			
CC	-!- FUNCTION: CATALYZES THE METHYLATION OF BOTH C-5 AND C-15 IN PRECORRIN-6Y TO FORM PRECORRIN-8X (BY SIMILARITY).			
CC	-!- CATALYTIC ACTIVITY: 2 S-adenosyl-L-methionine + precorrin-6Y = 2 S-adenosyl-L-homocysteine + precorrin-8X + CO(2).			
CC	-!- SIMILARITY: TO S-TYPEHIMURIUM CBIE; ALSO, LOW, TO OTHER METHYLASES INVOLVED IN COBALAMIN BIOSYNTHESIS.			

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EMBL: U67593; AAB99541.1; -  
PIR: A64490; A64490.  
TrEMBL: A64490; A64490.  
InterPro: IPR000878; Cor/por\_Mettransf.  
Pfam: PF00590; Tr\_methylase; 1.  
Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;  
Methyltransferase; Complete proteome.  
KW SEQUENCE 211 AA; 23805 MW; 279A1A2B14369510 CRC64;

Query Match 63.2%; Score 43; DB 1; Length 211;

Hypothetical protein; Lipid degradation; Hydrolase; Glycoprotein;  
Signal.

KW            1     19       POTENTIAL.  
KW SIGNAL  
FT CHAIN      20   633   LYSOPHOSPHOLIPASE C1450.09C.  
FF CARBOHYD   118   118   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   153   153   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   187   187   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   232   232   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   256   256   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   264   264   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   293   293   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   331   331   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   360   360   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   367   367   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   400   400   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   403   403   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   474   474   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   508   508   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   513   513   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   537   537   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   564   564   N-LINKED (GLNAC. .) (POTENTIAL).  
FT CARBOHYD   586   586   N-LINKED (GLNAC. .) (POTENTIAL).  
FF CARBOHYD   603   603   N-LINKED (GLNAC. .) (POTENTIAL).  
SQ SEQUENCE   633 AA; 68292 MW; 49871B29S5893DL9 CRC64;

Query Match          63.2%; Score 43; DB 1; Length 633;  
Best Local Similarity 69.2%; Pred. No. 11;  
Matches    9; Conservative    1; Mismatches    3; Indels    0; Gaps    0;

QY    1 PAVVGLSPGQEY 13  
||| ||| ||| |:  
DB    76 PASDGLSTGEQEF 88

RESULT 3  
RUVX CAUCR                   STANDARD;          PRF;      156 AA.  
ID - RUVX CAUCR           AC Q9A5K8;  
DT 10-OCT-2003 (Rel. 42, Created)  
DD 10-OCT-2003 (Rel. 42, Last sequence update)  
DE 10-OCT-2003 (Rel. 42, Last annotation update)  
DR Putative Holliday junction resolvase (EC 3.1.-.-.).  
GN C22439.

OS Caulobacter crescentus;  
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacterales;  
NC Oculobacteraceae; Caulobacter.  
OX NCBI\_TaxId=155892;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=ATCC 19089 / CBLS;  
RX MEDLINE=21173698; Pubmed=11259647;  
RA Niernman W.C., Feildilyum T.V., Laub M.T., Paulsen I.T., Nelson K.E.,  
RE Elsen J.A., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,  
RF Potocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,  
RG DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.L., Haft D.H.,  
RH Kolony J.F., Smit J., Craven M.B., Khouri H., Shetty J., Berry K.,  
RI Uterback T., Tran X., Wolf A., Vamathevan J., Ermolaeva M., White O.,  
RJ Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;  
RK "Complete genome sequence of *Caulobacter crescentus*.";  
RL Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001).

-!- FUNCTION: Could be a nuclease that resolves Holliday junction intermediates in genetic recombination.  
-!- SUBCELLULAR LOCATION: Cytoplasmic (potential).  
-!- SIMILARITY: Belongs to the YqgF HJR family.

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CC  EMBL; Z73966; CAA98214.1; -.
DR  EMBL; AE007063; AAK46406.1; -.
DR  EMBL; BX248341; CAD96945.1; -.
DR  PIR; E70764; E70764.
DR  TIGR; MT2126; -.
DR  TubercuList; RV2066; -.
DR  InterPro; IPR006364; Cobi_Cbil.
DR  InterPro; IPR006363; Cobi_Cbil.
DR  InterPro; IPR00878; Cor/por_Mettransf.
DR  InterPro; IPR003043; Uropor_Mettransf.
DR  Pfam; PF00590; TP_methylase; 2.
DR  TIGRfams; TIGR01467; cobi_cbil; 1.
DR  TIGRfams; TIGR01466; cobi_cbil; 1.
DR  PROSITE; PS00839; SUMT_1; 1.
DR  PROSITE; PS00840; SUMT_2; 1.
DR  KEGG; C06401; C06401; 1.
KW  Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW  Methyltransferase; Multifunctional enzyme; Complete proteome.
FT  DOMAIN 1 243 PRECORRIN-2 C20-METHYLTRANSFERASE.
FT  DOMAIN 244 508 PRECORRIN-3 METHYLASE.
FT  SEQUENCE 508 AA; 53910 MW; 95AC066F022C4DC1 CRC64;
SQ
Query Match 60.3%; Score 41; DB 1; Length 508;
Best Local Similarity 58.3%; Pred. No. 20;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY  2 AVVGLSPGEQY 13
DB  250 AVVGLGDSWD 261

RESULT 6
PLB4_SCHPO
ID  PLB4_SCHPO STANDARD; PRT; 673 AA.
AC  Q9P327;
DT  10-OCT-2003 (Rel. 42, Last sequence update)
DT  10-OCT-2003 (Rel. 42, Last sequence update)
DE  Putative lysophospholipase C977.09c precursor (EC 3.1.1.5)
DE  (Phospholipase B).
OS  SPAC977.09C OR SPAC1348.10C.
SN  Schizosaccharomyces pombe (Fission yeast).
OC  Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC  Schizosaccharomycetales; Schizosaccharomycetaceae;
OC  Schizosaccharomycetes.
OX  NCBI_TaxID=4896;
RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN=972;
RX  MEDLINE=2184401; PubMed=11859360;
RA  Wood V., Williams R., Rajadream M.A., Lyne M., Lyne R., Stewart A.,
RA  Sgouros J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA  Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA  Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,
RA  Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA  Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA  James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA  Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA  Oliver K., O'Neil S., Pearson D., Quail M.A., Rabinowitsch E.,
RA  Rutherford K., Rutter S., Saunders R., Seeger K., Sharp S.,
RA  Skelton J., Simmonds M., Squares S., Squares S., Stevens K.,
RA  Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA  Woodward J., Volckaert G., Aert R., Robben J., Grymonprez B.,
RA  Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA  Gabel C., Fuchs M., Fritzc C., Holzer E., Moestl D., Hilbert H.,
RA  Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Fohl T.M.,
RA  Eger P., Zimmermann W., Wedler H., Wambutt R., Furnelle B.,
RA  Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA  Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA  Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA  Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA  Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA  Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,

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RA  Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;
RT  "The genome sequence of Schizosaccharomyces pombe.";
RL  Nature 415:871-880(2002).
CC  -|- FUNCTION: Catalyzes the release of fatty acids from
CC  lysophospholipids (by similarity).
CC  -|- CATALYTIC ACTIVITY: 2-lysophosphatidylcholine + H(2)O =
CC  glycerophosphocholine + a fatty acid anion.
CC  -|- SUBCELLULAR LOCATION: Secreted (Probable).
CC  -|- SIMILARITY: Belongs to the lysophospholipase family.
CC
CC  This SWISS-PROT entry is copyright. It is produced through a collaboration
CC  between the Swiss Institute of Bioinformatics and the EMBL outstation
CC  the European Bioinformatics Institute. There are no restrictions on its
CC  use by non-profit institutions as long as its content is in no way
CC  modified and this statement is not removed. Usage by and for commercial
CC  entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC  or send an email to license@isb-sib.ch).
CC
CC  EMBL; AL358912; CAB94277.1; -.
CC  EMBL; AL37130; CAB9631.1; -.
CC  PIR; T50281; T50281.
CC  GenDB SPombe; SPAC977.09c; -.
CC  InterPro; IPR001179; SKBP_PPIase.
CC  InterPro; IPR002642; PLAC.
CC  Pfam; PF01735; PLA2_B; 1.
CC  SMART; SMC0022; PLAC; 1.
CC  KW  Hypothetical protein; Lipid degradation; Hydrolase; Glycoprotein;
CC  Signal.
CC  FT  SIGNAL 1 19 POTENTIAL.
CC  FT  CHAIN 20 673 PUTATIVE LYSOPHOSPHOLIPASE C977.09C.
CC  FT  CARBOHYD 72 72 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 125 125 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 191 191 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 194 194 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 272 272 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 301 301 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 374 374 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 404 404 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 409 409 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 481 481 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 516 516 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 545 545 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  FT  CARBOHYD 574 574 N-LINKED (GLCNAC. .) (POTENTIAL).
CC  SQ  SEQUENCE 673 AA; 74595 MW; B39A77E76CD694B CRC64;
Query Match 58.8%; Score 40; DB 1; Length 673;
Best Local Similarity 61.5%; Pred. No. 39;
Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY  1 PAVVGLSPGEQY 13
DB  83 PASEGLNEGEQSY 95

RESULT 7
KV31_HUMAN
ID  KV31_HUMAN STANDARD; PRT; 115 AA.
AC  P04433;
DT  13-AUG-1987 (Rel. 05, Created)
DT  13-AUG-1987 (Rel. 05, Last sequence update)
DT  15-JUL-1999 (Rel. 38, Last annotation update)
DE  Ig kappa chain V-III region VG precursor (Fragment).
OS  Homo sapiens (Human).
OC  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX  NCBI_TaxID=9606;
RN  [1]
RP  SEQUENCE FROM N.A.
RC  MEDLINE=85087932; PubMed=6440122;
RX  Pech M., Zachau H.G.;
RA  "Immunoglobulin genes of different subgroups are interdigitated
RT  within the VK locus";
RL  Nucleic Acids Res. 12:9229-9236(1984).

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EMBL; X01668; -; NOT_ANNOTATED_CDS.
DR PIR; A01900; K3HUVG.
DR HSP; P80362; 1MTL.
DR GO; GO:0005576; C:extracellular; NAS.
DR GO; GO:0003823; F:antigen binding; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; IG-like.
DR InterPro; IPR003596; IG_V.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS0835; IG LIKE; 1.
KW Immunoglobulin V region; Signal.
FT SIGNAL 1 20
FT CHAIN 21 >115 IG KAPPA CHAIN V-III REGION VG.
FT DOMAIN 21 43 FRAMEWORK-1.
FT DOMAIN 44 54 COMPLEMENTARITY-DETERMINING-1.
FT DOMAIN 55 69 FRAMEWORK-2.
FT DOMAIN 70 76 COMPLEMENTARITY-DETERMINING-2.
FT DOMAIN 77 108 FRAMEWORK-3.
FT DOMAIN 109 115 COMPLEMENTARITY-DETERMINING-3.
FT DISULFID 43 108 BY SIMILARITY.
FT NON TER 115 115
FT SEQUENCE 115 AA; 12575 MW; 2DE47CDA3A17D555 CRC64;
Query Match 57.4%; Score 39; DB 1; Length 115;
Best Local Similarity 63.6%; Pred. No. 10;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
QY 1 PAVVGLSPGEQ 11
DB 28 PATLSVSPGER 38
RESULT 8
KVSI_MOUSE
ID KVSI_MOUSE STANDARD; PRT; 115 AA.
AC P01642;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Ig kappa chain V-V region L7 precursor (Fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=81220975; PubMed=6264318;
RA Pech M., Hochtl J., Schnell H., Zachau H.G.;
RT "Differences between germ-line and rearranged immunoglobulin V kappa
coding sequences suggest a localized mutation mechanism."
RL Nature 291:668-670(1981).
CC -1- MISCELLANEOUS: THERE APPEAR TO BE TWO POSSIBLE SPLICE JUNCTIONS AT
CC THE 3' END OF THE INTRON. THE ALTERNATE WOULD CODE FOR A PROTEIN
CC LACKING RESIDUES 17-19.
DR PIR; A01925; KVM5L7.
DR PDB; 1J10; 18-FEB-03.
DR PDB; 1J1P; 18-FEB-03.
DR PDB; 1J1X; 18-FEB-03.
DR InterPro; IPR007110; IG-like.
DR InterPro; IPR003596; IGV.
DR Pfam; PF00047; IGV; 1.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS0835; IG LIKE; 1.

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KW Immunoglobulin V region; Signal; 3D-structure.
FT SIGNAL 1 20
FT CHAIN 21 >115 IG KAPPA CHAIN V-V REGION L7.
FT DOMAIN 21 43 FRAMEWORK-1.
FT DOMAIN 44 54 COMPLEMENTARITY-DETERMINING-1.
FT DOMAIN 55 69 FRAMEWORK-2.
FT DOMAIN 70 76 COMPLEMENTARITY-DETERMINING-2.
FT DOMAIN 77 108 FRAMEWORK-3.
FT DOMAIN 109 >115 COMPLEMENTARITY-DETERMINING-3.
FT DISULFID 43 108 BY SIMILARITY.
FT NON TER 115 115
FT SEQUENCE 115 AA; 12615 MW; C17BEC758C577E00 CRC64;
Query Match 57.4%; Score 39; DB 1; Length 115;
Best Local Similarity 54.5%; Pred. No. 10;
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
QY 1 PAVVGLSPGEQ 11
DB 28 PATLSVSPGER 38
RESULT 9
BUD3_YEAST
ID BUD3_YEAST STANDARD; PRT; 1636 AA.
AC P25558; P25556; P25557; P87007;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Bud site selection protein BUD3.
DE BUD3 OR YCL014W/YCL013W/YCL012W OR YCL14W/YCL13W/YCL12W.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=95247824; PubMed=7730410;
RA Chant J., Mischke M., Mitchell E., Herskowitz I., Pringle J.R.;
RT "Role of Bud3p in producing the axial budding pattern of yeast."
RL J. Cell Biol. 129:767-778(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=S288c;
RX MEDLINE=92244356; PubMed=1574125;
RA Oliver S.G., van der Aart Q.J.M., Agostoni-Carbone M.L., Aigle M.,
RA Alberghina L., Alexandraki D., Antoine G., Anwar R., Bailesta J.P.G.,
RA Benit P., Berben G., Bergantino E., Biteau N., Bolle P.-A.,
RA Bolotin-Fukuhara M., Brown A.J.P., Brown R., Buhler J.-M.,
RA Carignani G., Chanet R., Contreras R., Crouzet M., Daignan-Fornier B.,
RA De Haan M., Defoor E., Delgado M.D., Demolder J., Doira C., Dubois E.,
RA Dujon B., Duesterhoeft A., Erdmann H., Fiers W.,
RA Fairhead C.A., Faye G., Feldmann L., Frontali L., Fukuhara H.,
RA Frangoules-Gaillard M.-C., Franco L., Gansdorff N.,
RA Fuller L.J., Gent M.E., Gigot D., Gilliquet V., Glandsdorff N.,
RA Goffeau A., Grenson M., Grisanti P., Grivell L.A., Haasemann M.,
RA Hatat D., Hegemann J.H., Herbert C.J., Hilger F., Hohmann S.,
RA Hollenberg C.P., Huse K., Iborra F., Indge K.J., Isono K., Jackman P.,
RA Jacq C., Jaquet M., James C.M., Jauniaux J.-C., Jia Y., Jimenez A.,
RA Kleinhans U., Kreis P., Lafranchi G., Lewis C., van der Linden C.G.,
RA Lucchini G., Luttenkirchen K., Maat C., Mannheim G., Manzano M.E.,
RA Martegani E., Mathieu A., Maurer C.T.C., McConnell D., McKee R.A.,
RA Messenguy F., Mewes H.-W., Molemans F., Montague M.A., Navas L.,
RA Newlon C.S., Olson M.V., Pallier C., Panzeri L., Pearson B.M.,
RA Perea J., Philippsen P., Pierard A., Planta R.J., Plevani S.W.,
RA Poetsch B., Pohl F.M., Purnelle B., Ramezani Rad M., Rasmussen S.W.,
RA Raynal A., Remacha M., Richterich P., Roberts A.B., Rodriguez F.,
RA Sanz E., Schaaff-Gerstenschlaeger I., Scherens B., Schweitzer B.,
RA Shu Y., Skala J., Slonimski P.P., Sor F., Soustelle C.,
RA Spiegelberg R., Staveva L.I., Steensma H.Y., Steiner S., Thierry A.,
RA Thireos G., Triano L.N., Urrestazu L.A., Valle G., Vetter I.,
RA van Vliet-Reedijk J.C., Volckaert G., Vreken P., Warmington J.R.,
RA von Wettstein D., Wickstead B.L., Wilson C., Wurst H., Xu G.,

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RA Zimmermann F.K., Sgouros J.G.;
RT "The complete DNA sequence of yeast chromosome III.";
RL Nature 357:38-46(1992).
[3]
RP REVISIONS.
RA Gromadka R.;
RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
[4]
RN REVISIONS.
RA Valles G., Volckaerts G.;
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Co-assembles with BUD4 at bud sites. BUD4 and BUD3 may
CC cooperate to recognize a spatial landmark (the neck filaments)
CC during mitosis and they subsequently become a landmark for
CC establishing the axial budding pattern in G1.
CC -----
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CC -----
DR EMBL; U17580; AAA86315.1; -.
DR EMBL; X59720; CA42346.2; -.
DR PIR; S74285; S74285.
DR PIR; S74286; S74286.
DR GeneOnline; 138855; -.
DR SGD; S0000520; BUD3.
DR InterPro; IPR000219; RhoGEF.
DR SMART; SM00325; RhoGEF; 1.
KW Cell cycle.
SQ SEQUENCE 1636 AA; 184717 MW; 9E4E46BA5C3A3F69 CRC64;

Query Match 57.4%; Score 39; DB 1; Length 1636;
Best Local Similarity 58.3%; Pred.No. 1.4e+02;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 12
||: ||| |||
DB 112 PAIENLSPDQE 123

RESULT 10
RS1 LEULA STANDARD; PRT; 429 AA.
AC P50889; P71450;
DT 01-OCT-1996 (Rel. 34, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE 40S ribosomal protein S1.
GN RPS1.
OS Leuconostoc lactis.
OC Bacteria; Firmicutes; Lactobacillales; Leuconostoc.
OX NCBI_TaxID=1246;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97186703; PubMed=9034319;
RA Yamit-Rezi A., Levy Z., Neuman S., Nudel U.;
RT "A Leuconostoc lactis protein with homology to ribosomal protein S1
RT shares common epitopes and common DNA binding properties with a
RT mammalian DNA binding nuclear factor.";
RL Gene 185:99-103(1997).
[2]
RN [2]
RP SEQUENCE OF 24-429 FROM N.A.
RX MEDLINE=95237615; PubMed=7721096;
RA Eklund E.A., Lee S.W., Skalniak D.G.;
RT "Cloning of a cDNA encoding a human DNA-binding protein similar to
RT ribosomal protein S1.";
RL Gene 155:231-235(1995).
[3]
RN [3]
RP SEQUENCE OF 78-429 FROM N.A.

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RX MEDLINE=96164600; PubMed=8568274;
RA Tuzaka K., Ieu A.K., Frank M.B., Movafagh B.F., Koscec M.,
RA Winkler T.H., Kalden J.R., Reichlin M.;
RT "Lupus autoantibodies to double-stranded DNA cross-react with
RT ribosomal protein S1.";
RL J. Immunol. 156:1668-1675(1996).
CC -!- FUNCTION: EXHIBITS PREFERENTIAL BINDING TO SINGLE-STRANDED AND
CC DOUBLE-STRANDED DNA AND A LOW BINDING AFFINITY FOR RNA.
CC -!- SIMILARITY: Belongs to the S1P family of ribosomal proteins.
CC -!- SIMILARITY: Contains 4 S1 motif domains.
CC -!- CAUTION: WAS ORIGINALLY (REF.2 AND REF.3) THOUGHT TO ORIGINATE
CC FROM HUMAN BUT IS MOST PROBABLY THE RESULT OF A CDNA LIBRARY
CC CONTAMINATION BY L.LACTIS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; U24086; AAB08978.1; -.
DR EMBL; U05589; AAA77669.1; -.
DR EMBL; U27517; AAA97575.1; -.
DR HSP; P05055; ISRO.
DR InterPro; IPR008994; Nucleic_acid_OB.
DR InterPro; IPR00110; Ribosomal_S1.
DR InterPro; IPR003029; S1.
DR Pfam; PF00575; S1; 4.
DR PRINTS; PR00681; RIBOSOMALS1.
DR SMART; SM00316; S1; 4.
DR PROSITE; PS50126; S1; 4.
KW Ribosomal protein; Repeat; RNA-binding.
FT DOMAIN 55 128 S1 MOTIF 1.
FT DOMAIN 144 211 S1 MOTIF 2.
FT DOMAIN 231 299 S1 MOTIF 3.
FT DOMAIN 316 385 S1 MOTIF 4.
FT CONFLICT 24 24 S -> G (IN REF. 2).
FT CONFLICT 122 122 A -> S (IN REF. 3).
FT CONFLICT 217 217 L -> R (IN REF. 2 AND 3).
SQ SEQUENCE 429 AA; 46386 MW; 92AC82605F39DDFC CRC64;

Query Match 55.9%; Score 38; DB 1; Length 429;
Best Local Similarity 80.0%; Pred.No. 55;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 AVVGLSPGEQ 11
||||| |||
DB 71 AVVGLSTGEE 80

RESULT 11
CATA_PICAN STANDARD; PRT; 507 AA.
ID CATA_PICAN STANDARD; PRT; 507 AA.
AC P30263;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Peroxisomal catalase (BC 1.11.1.6).
GN PXP9 OR PXP-9.
OS Pichia angusta (Yeast) (Hansenula polymorpha).
OC Eukaryota; Fungi; Ascomycota; Saccharomycetina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Pichia.
OX NCBI_TaxID=4905;
RN [1]
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=ATCC 34438;
RA MEDLINE=92299073; PubMed=1607006;
RA Didion T., Roggenkamp R.O.;
RT "Targeting signal of the peroxisomal catalase in the methylotrophic
RT yeast Hansenula polymorpha.";
RL FEBS Lett. 303:113-116(1992).

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CC -!- FUNCTION: Occurs in almost all aerobically respiring organisms and  
 CC serves to protect cells from the toxic effects of hydrogen  
 CC peroxide.  
 CC -!- CATALYTIC ACTIVITY: 2 H(2)O(2) = O(2) + 2 H(2)O.  
 CC -!- COFACTOR: Heme group.  
 CC -!- SUBUNIT: Homotetramer.  
 CC -!- SUBCELLULAR LOCATION: Peroxisomal.  
 CC -!- SIMILARITY: Belongs to the catalase family.  
 CC  
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 CC  
 CC EMBL; X56501; CRA39856.1; -.  
 CC PIR; S23422; S23422.  
 CC HSP; P15202; IA4E.  
 CC InterPro; IPR002226; Catalase.  
 CC Pfam; PF001199; Catalase; 1.  
 CC PRINTS; PR00067; CATALASE.  
 CC ProDom; PD000510; Catalase; 1.  
 CC PROSITE; PS00437; CATALASE 1; 1.  
 CC PROSITE; PS00438; CATALASE 2; 1.  
 CC Oxidoreductase; Peroxidase; Iron; Heme; Hydrogen peroxide;  
 CC Peroxisome.  
 CC ACT\_SITE 65 65 BY SIMILARITY.  
 CC FT ACT\_SITE 138 138 BY SIMILARITY.  
 CC FT METAL 348 348 IRON (HEME AXIAL LIGAND) (BY SIMILARITY).  
 CC FT SITE 505 507 MICROBODY TARGETING SIGNAL (POTENTIAL).  
 CC SEQUENCE 507 AA; 57849 MW; 3536ED0A49539CC3 CRC64;  
 CC  
 CC Query Match 55.9%; Score 38; DB 1; Length 507;  
 CC Best Local Similarity 70.0%; Pred. No. 65;  
 CC Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
 CC  
 CC QY 3 VVGLSPGEQE 12  
 CC | : : : : :  
 CC Db 445 VLGRTPGEEQ 454  
 CC  
 CC RESULT 12  
 CC COBJ\_ARCFU STANDARD; PRT; 446 AA.  
 CC ID\_COBJ\_ARCFU STANDARD; PRT; 446 AA.  
 CC AC O29534;  
 CC DT 15-JUL-1998 (Rel. 36, Created)  
 CC DT 15-JUL-1998 (Rel. 36, Last sequence update)  
 CC DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 CC DE Cobalamin biosynthesis protein cobJH [includes: Precorrin-3B C17-  
 CC methyltransferase [EC 2.1.1.131] (Precorrin-3 methyltransferase)  
 CC DE (Precorrin-3 methylase); Precorrin-8X methylmutase (EC 5.4.1.2)  
 CC DE (Precorrin isomerase)].  
 CC COBJH OR AF0724.  
 CC GN COBJH  
 CC OS Archaeoglobus fulgidus  
 CC OC Archaea; Euryarchaeota; Archaeoglobi; Archaeoglobales;  
 CC OC Archaeoglobaceae; Archaeoglobus.  
 CC OX NCBI\_TaxID=2234;  
 CC [1]  
 CC SEQUENCE FROM N.A.  
 CC RC STRAIN=VC-16 / DSM 4304 / ATCC 49558;  
 CC RX MEDLINE=98049343; PubMed=989475;  
 CC RA Klenk H.-P., Clayton R.A., Tomb J.-F., White O., Nelson K.E.,  
 CC Ketchum K.A., Dodson R.J., Gwinn M., Hickey E.K., Peterson J.D.,  
 CC Richardson D.L., Kerlavage A.R., Graham D.B., Kyripides N.C.,  
 CC Fleischmann D., Quackenbush J., Lee N.H., Sutton G.G., Gill S.,  
 CC Kirkness E.F., Dougherty B.A., McKenney K., Adams M.D., Loftus B.,  
 CC Peterson S., Reich C.I., McNeil L.K., Badger J.H., Glodek A., Zhou L.,  
 CC Overbeek R., Gayney J.D., Weidman J.F., McDonald L., Uterback T.,  
 CC Cotton M.D., Spriggs T., Artlach P., Kaine B.P., Sykes S.M.,  
 CC Sadow P.W., D'Andrea K.P., Bowman C., Fujii C., Garland S.A.,  
 CC Mason T.M., Olsen G.J., Fraser C.M., Smith H.O., Woese C.R.,

RA Venter J.C.;  
 RT "The complete genome sequence of the hyperthermophilic, sulphate-  
 RL reducing archaeon *Archaeoglobus fulgidus*.";  
 CC Nature 390:364-370(1997).  
 CC -!- FUNCTION: BIFUNCTIONAL ENZYME WITH A METHYLTRANSFERASE DOMAIN THAT  
 CC CATALYZES THE METHYLATION OF C-17 IN PRECORRIN-3B TO FORM  
 CC PRECORRIN-4 AND AN ISOMERASE DOMAIN THAT CATALYZES THE CONVERSION  
 CC OF PRECORRIN-8X TO HYDROGENOBYRINIC ACID; A METHYL MIGRATION  
 CC REACTION DURING THE TRANSFORMATION OF PRECORRIN-3 TO FORM  
 CC COBYRINIC ACID (BY SIMILARITY).  
 CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + precorrin-3B = S-  
 CC adenosyl-L-homocysteine + precorrin 4.  
 CC -!- CATALYTIC ACTIVITY: Precorrin-8X = hydrogenobyrinate.  
 CC -!- PATHWAY: Cobalamin biosynthesis.  
 CC -!- SIMILARITY: Belongs to the cobH family.  
 CC  
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 CC  
 CC EMBL; AE001055; AAB90518.1; -.  
 CC PIR; D69340; D69340.  
 CC TIGR; AF0724; -.  
 CC InterPro; IPR003722; CbiC.  
 CC InterPro; IPR006363; CobJ.  
 CC InterPro; IPR000878; Cor/por\_Metransf.  
 CC Pfam; PF02570; CbiC; 1.  
 CC Pfam; PF00590; TP\_methylase; 1.  
 CC TIGRFAMS; TIGR01456; cobJ cbiH; 1.  
 CC Cobalamin biosynthesis; Transferase; Methyltransferase; Isomerase;  
 CC KW Multifunctional enzyme; Complete proteome.  
 CC FT DOMAIN 1 246 PRECORRIN-3B C-17 METHYLTRANSFERASE.  
 CC FT DOMAIN 247 446 PRECORRIN-8X METHYLMUTASE.  
 CC SEQUENCE 446 AA; 48678 MW; 7341BCBC0998FFDA CRC64;  
 CC  
 CC Query Match 54.4%; Score 37; DB 1; Length 446;  
 CC Best Local Similarity 60.0%; Pred. No. 85;  
 CC Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;  
 CC  
 CC QY 3 VVGLSPGEQE 12  
 CC | : : : : :  
 CC Db 12 VVGIGPKEE 21  
 CC  
 CC RESULT 13  
 CC ENOI\_MAIZE STANDARD; PRT; 446 AA.  
 CC ID\_ENOI\_MAIZE STANDARD; PRT; 446 AA.  
 CC AC P26301; 1992 (Rel. 22, Created)  
 CC DT 01-MAY-1992 (Rel. 22, Last sequence update)  
 CC DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 CC DE Enolase 1 (EC 4.2.1.11) (2-phosphoglycerate dehydratase 1) (2-phospho-  
 CC D-glycerate hydro-lyase 1).  
 CC GN ENOI OR PGH1.  
 CC OS Zea mays (Maize).  
 CC OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 CC OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
 CC OC PACCAD clade; Panicoideae; Andropogoneae; Zea.  
 CC OX NCBI\_TaxID=4577;  
 CC [1]  
 CC SEQUENCE FROM N.A.  
 CC RC STRAIN=cv. Berkeley Fast; TISSUE=Root;  
 CC RX MEDLINE=91316216; PubMed=1859865;  
 CC RA Lal S.K., Johnson S., Conway T., Kelley P.M.;  
 CC "Characterization of a maize cDNA that complements an enolase-  
 CC deficient mutant of *Escherichia coli*.";  
 CC Plant Mol. Biol. 16:787-795(1991).  
 CC -!- CATALYTIC ACTIVITY: 2-phospho-D-glycerate = phosphoenolpyruvate +  
 CC H(2)O.

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CC -|- COFACTOR: Magnesium is required for catalysis and for stabilizing
CC the dimer (by similarity).
CC -|- PATHWAY: Glycolysis.
CC -|- SUBUNIT: Homodimer.
CC -|- SIMILARITY: Belongs to the enolase family.
CC -----
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CC -----
DR ENBL; X55981; CAA39454.1; -.
DR PIR; S16257; S16257.
DR HSP; P56252; 1PZD.
DR MaizedB; 30060; -.
DR InterPro; IPR000941; Enolase.
DR Pfam; PF00113; enolase_1.
DR Pfam; PF03952; enolase_N; 1.
DR PRINTS; PR00148; ENOLASE.
DR ProDom; PD000902; eno; 1.
DR TIGRFAMs; TIGR01060; eno; 1.
DR PROSITE; PS00164; ENOLASE; 1.
DR Lyase; Glycolysis; Magnesium; Multigene family.
FT ACT SITE 164 164 BY SIMILARITY.
FT METAL 251 251 MAGNESIUM (BY SIMILARITY).
FT METAL 302 302 MAGNESIUM (BY SIMILARITY).
FT METAL 329 329 MAGNESIUM (BY SIMILARITY).
SQ SEQUENCE 446 AA; 48063 MW; 6266C48914F35198 CRC64;

Query Match 54.48; Score 37; DB 1; Length 446;
Best Local Similarity 63.63; Pred. No. 85;
Matches 7; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 11
   |||||
Db 77 PAIVGKDPTQ 87

RESULT 14
NR41 RAT STANDARD; PRT; 597 AA.
AC P22829;
DT 01-AUG-1991 (Rel. 19, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Orphan nuclear receptor HMR (Nerve growth factor induced protein I-B)
DE (NGFI-B) (NUR77).
GN NR4A1 OR HMR OR NGFI-B.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90166506; PubMed=3272167;
RA Milbrandt J.;
RT "Nerve growth factor induces a gene homologous to the glucocorticoid
RT receptor gene."
RL Neuron 11:183-188 (1988).
RN [2]
RP CHARACTERIZATION.
RX MEDLINE=93361012; PubMed=8395013;
RA Wilson T.E., Fahrner T.J., Milbrandt J.;
RT "The orphan receptors NGFI-B and steroidogenic factor 1 establish
RT monomer binding as a third paradigm of nuclear receptor-DNA
RT interaction."
RL Mol. Cell. Biol. 13:5794-5804 (1993).
RN [3]
RP DNA BINDING MOTIFS.

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RX MEDLINE=92229411; PubMed=1314418;
RA Wilson T.E., Paulsen R.E., Padgett K.A., Milbrandt J.;
RT "Participation of non-zinc finger residues in DNA binding by two
RT nuclear orphan receptors."
RL Science 256:1107-1110 (1992).
RN [4]
RP PHOSPHORYLATION.
RX MEDLINE=94043340; PubMed=8227042;
RA Hirata Y., Kiuchi K., Chen H.-C., Milbrandt J., Guroff G.;
RT "The phosphorylation and DNA binding of the DNA-binding domain of the
RT orphan nuclear receptor NGFI-B."
RL J. Biol. Chem. 268:24808-24812 (1993).
RN [5]
RP X-RAY CRYSTALLOGRAPHY (2.7 ANGSTROMS) OF 264-351 IN COMPLEX WITH NBRE,
RP AND METAL-BINDING.
RX MEDLINE=99260743; PubMed=10331876;
RA Meinke G., Sigler P.B.;
RT "DNA-binding mechanism of the monomeric orphan nuclear receptor
RT NGFI-B."
RL Nat. Struct. Biol. 6:471-477 (1999).
CC -|- FUNCTION: Probable nuclear receptor. May act concomitantly with
CC NUR1 in regulating the expression of delayed-early genes during
CC liver regeneration. Binds the NGFI-B response element (NBRE) 5'-
CC AAAAGGTCA-3'.
CC -|- SUBUNIT: Binds DNA as a monomer.
CC -|- SUBCELLULAR LOCATION: Nuclear.
CC -|- TISSUE SPECIFICITY: Expressed in lung, brain and superior
CC cervical ganglia. High levels are seen in the adrenal tissue.
CC -|- INDUCTION: By nerve growth factor and during liver regeneration.
CC -|- PTM: Phosphorylation of Ser-350 results in decrease in NBRE
CC binding while phosphorylation of Ser-340 has little effect on it.
CC -|- SIMILARITY: Belongs to the nuclear hormone receptor family. NR4
CC subfamily.
CC -----
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CC -----
DR ENBL; U17254; AAA56770.1; ALT_INIT.
DR PDB; 1CIT; 26-JUN-00.
DR TRAFAC; T00619; -.
DR InterPro; IPR000536; Hormone rec lig.
DR InterPro; IPR001723; Steroid receptor.
DR InterPro; IPR008946; Str_ncl_receptor.
DR InterPro; IPR001628; Znf_C4steroid.
DR Pfam; PF00104; hormone_rec; 1.
DR Pfam; PF00105; zf-C4; 1.
DR PRINTS; PR00398; STRDHORMNER.
DR PRINTS; PR00047; STROIDFINGER.
DR ProDom; PD000035; Znf_C4steroid; 1.
DR SMART; SM00430; HOL1; 1.
DR SMART; SM00399; Znf_C4; 1.
DR PROSITE; PS00031; NUCLEAR RECEPTOR; 1.
KW Receptor; Transcription regulation; DNA-binding; Nuclear protein;
KW Zinc-finger; Phosphorylation; 3D-structure.
FT DNA_BIND 266 331 NUCLEAR RECEPTOR-TYPE.
FT ZN_FING 266 286 C4-TYPE.
FT ZN_FING 302 326 C4-TYPE.
FT DOMAIN 408 458 LIGAND-BINDING (POTENTIAL).
FT DOMAIN 80 91 POLY-SER.
FT DOMAIN 182 186 POLY-PRO.
FT DOMAIN 582 585 POLY-PRO.
FT MOD_RES 340 340 PHOSPHORYLATION (BY PKA).
FT MOD_RES 350 350 PHOSPHORYLATION (BY PKA).
FT MUTAGEN 340 340 S->A: LOSS OF PHOSPHORYLATION.
FT MUTAGEN 350 350 S->A: LOSS OF PHOSPHORYLATION.
FT MUTAGEN 345 345 R->K: DECREASED NBRE BINDING.
FT MUTAGEN 348 348 L->V: ALMOST COMPLETE LOSS OF NBRE
FT BINDING.

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SQ SEQUENCE 597 AA; 64281 MW; 9CFA987112337E53 CRC64;
Query Match 54.4%; Score 37; DB 1; Length 597;
Best Local Similarity 50.0%; Pred. No. 1.1e+02;
Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQE 12
| : |||||:
Db 424 PGFELSFGPD 435

RESULT 15
GLMS CLOTE STANDARD; PRT; 607 AA.
AC Q890U2;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Glucosamine--fructose-6-phosphate aminotransferase [isomerizing]
DE (EC 2.6.1.16) (Hexosephosphate aminotransferase) (D-fructose-6-
DE phosphate amidotransferase) (GFAT) (L-glutamine-D-fructose-6-phosphate
DE amidotransferase) (Glucosamine-6-phosphate synthase).
GN GLMS OR C1C02543.
OS Clostridium tetani.
OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
OC Clostridium.
OX NCBI_TaxID=1513;
RN [1]
SEQUENCE FROM N.A.
RP STRAIN=Massachusetts / E88;
RX MEDLINE=22457253; PubMed=12552129;
RA Brueggemann H., Baumer S., Fricke W.F., Wierer A., Liesegang H.,
RA Decker I., Herzberg C., Martinez-Arias R., Merkl R., Henne A.,
RA Gottschalk G.;
RT "The genome sequence of Clostridium tetani, the causative agent of
RT tetanus disease.";
RL Proc. Natl. Acad. Sci. U.S.A. 100:1316-1321(2003).
CC -!- FUNCTION: Catalyzes the first step in hexosamine metabolism,
CC converting fructose-6P into glucosamine-6P using glutamine as a
CC nitrogen source (By similarity).
CC -!- CATALYTIC ACTIVITY: L-glutamine + D-fructose 6-phosphate = L-
CC Glutamate + D-glucosamine 6-phosphate.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: In the C-terminal section; belongs to the SIS family.
CC GFAT subfamily.
CC -!- SIMILARITY: Contains 1 type-2 glutamine amidotransferase domain.
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-----
DR EMBL; A0015944; AAC37003.1; ALT_INIT.
DR HAMAP; MF 00164; -; 1.
DR InterPro; IPR000583; GATase_2.
DR InterPro; IPR005855; GLMS.
DR InterPro; IPR001347; SIS.
DR Pfam; PF003110; GATase_2; 1.
DR Pfam; PF01380; SIS; 2.
DR TIGRPFAMs; TIGR01135; qlms; 1.
DR PROSITE; PS00443; GATASE_TYPE_II; 1.
KW Transferase; Aminotransferase; Glutamine amidotransferase;
KW Complete proteome.
FT INIT MET 0 0 BY SIMILARITY.
FT DOMAIN 1 239 GLUTAMINE AMIDOTRANSFERASE.
FT ACT_SITE 1 1 GATASE (BY SIMILARITY).
FT ACT_SITE 602 602 ISOMERIZATION FRU-6P (BY SIMILARITY).
SQ SEQUENCE 607 AA; 67694 MW; F4B1CFA2EEA37948 CRC64;

Query Match 54.4%; Score 37; DB 1; Length 607;
```

Best Local Similarity 61.5%; Pred. No. 1.1e+02;  
Matches 8; Conservative 2; Mismatches 1; Indels 2; Gaps 1;

QY 1 PAVVGLSPGEQEY 13  
| : |||||: ||:  
Db 176 PLIVGL--GEEY 186

Search completed: May 7, 2004, 12:34:33  
Job time : 4.68 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:27:20 ; Search time 8.32 Seconds  
(without alignments)  
150.299 Million cell updates/sec

Title: US-09-786-214A-13

Perfect score: 68  
Sequence: 1 PAVGLSPGEQY 13

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : PIR 78:\*

1: PIR1: \*  
2: PIR2: \*  
3: PIR3: \*  
4: PIR4: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	63.2	211	2 A64490	precorrin-6Y methy
2	43	63.2	623	2 T40991	probable lysophosp
3	43	63.2	821	2 C84304	DNA helicase limpo
4	41	60.3	156	2 F87551	conserved hypotet
5	41	60.3	326	2 T45226	probable N5,N10-me
6	41	60.3	508	2 E70764	probable cobI prote
7	40	58.8	227	2 B90400	hypothetical prote
8	40	58.8	243	2 I54459	MHC H-2K1-k - mous
9	40	58.8	381	2 F75270	cytochrome p450 -
10	40	58.8	540	2 A75250	carboxylesterase,
11	40	58.8	661	2 G84511	hypothetical prote
12	40	58.8	673	2 T50281	probable lysophosp
13	39	57.4	43	2 S21065	Ig kappa chain V r
14	39	57.4	96	2 S45441	Ig kappa chain V r
15	39	57.4	103	2 S19975	Ig kappa chain V r
16	39	57.4	106	2 PS0070	Ig kappa chain V r
17	39	57.4	106	2 PC4282	Ig kappa chain (an
18	39	57.4	107	2 S57444	Ig kappa chain V-J
19	39	57.4	108	2 C30502	Ig kappa chain V r
20	39	57.4	108	2 G44151	Ig kappa chain V r
21	39	57.4	111	2 S23628	Ig kappa chain V r
22	39	57.4	114	2 S54905	Ig kappa chain V r
23	39	57.4	115	1 K3HUVG	Ig kappa chain pre
24	39	57.4	115	1 KYMSL7	Ig kappa chain pre
25	39	57.4	115	2 S11697	Ig kappa chain pre
26	39	57.4	116	2 B25521	Ig kappa chain pre
27	39	57.4	119	2 S41816	Ig kappa chain V r
28	39	57.4	125	2 S40344	Ig kappa chain V-J
29	39	57.4	128	2 PN0445	Ig kappa chain pre

30 39 57.4 128 2 S40379 Ig kappa chain V-J  
31 39 57.4 128 2 A56701 Ig kappa chain V r  
32 39 57.4 129 2 S29627 Ig kappa chain V r  
33 39 57.4 129 2 S40363 Ig kappa chain - h  
34 39 57.4 132 2 S05268 Ig kappa chain pre  
35 39 57.4 144 2 P01006 Ig kappa chain pre  
36 39 57.4 144 2 P01006 Ig heavy chain V r  
37 39 57.4 215 2 A23746 Ig kappa chain V-I  
38 39 57.4 428 2 AG1304 uracil permease ho  
39 39 57.4 428 2 AG1676 uracil permease ho  
40 39 57.4 1367 2 S74285 BUD3 protein - yea  
41 38 55.9 86 2 S26649 Ig heavy chain V r  
42 38 55.9 110 2 S60591 Ig light chain var  
43 38 55.9 111 2 PN0537 Ig kappa chain V r  
44 38 55.9 152 2 G75184 probable transcrip  
45 38 55.9 155 2 G71217 probable transcrip

## ALIGNMENTS

RESULT 1  
A64490  
precorrin-6Y methylase homolog - Methanococcus jannaschii  
C:Species: Methanococcus jannaschii  
C:Date: 13-Sep-1996 #sequence\_revision 13-Sep-1996 #text\_change 18-Aug-2003  
C:Accession: A64490  
R:Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann, R.D.; Sutton, G.G.; Blake,  
; Reich, C.I.; Overbeek, R.; Kirkness, B.F.; Weinstock, K.G.; Merrick, J.M.; Glodek, A.,  
rson, J.D.; Sadow, P.W.; Hanna, M.C.; Cotton, M.D.; Roberts, K.M.; Hurst, M.A.  
Science 273, 1058-1073, 1996  
A:Authors: Kaine, B.P.; Borodovsky, M.; Klenk, H.P.; Fraser, C.M.; Smith, H.O.; Woese, C  
A:Title: Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii  
A:Reference number: A64300; MUID:96337999; PMID:8688087  
A:Accession: A64490  
A>Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Molecule type: DNA  
A:Residues: 1-211 <BUL>  
A:Cross-references: GB:U67593; GB:L77117; NID:g2826427; PIDN:AAB99541.1; PID:g1592152;  
C:Genetics:  
A:Map position: FOR1500322-1500957  
C:Superfamily: precorrin-6Y methylase ChiE

Query Match 63.2%; Score 43; DB 2; Length 211;  
Best Local Similarity 54.5%; Pred. No. 6.7;  
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 3 VVGLSPGEQY 13  
DB 4 IVGIGPGDREY 14

## RESULT 2

T40991  
probable lysophospholipase precursor - fission yeast (Schizosaccharomyces pombe)  
C:Species: Schizosaccharomyces pombe  
C:Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 21-Jan-2000  
C:Accession: T40991  
R:Lyne, M.; Rajandream, M.A.; Barrell, B.G.; Volckaert, G.  
submitted to the EMBL Data Library, March 1999  
A:Reference number: Z21962  
A:Accession: T40991  
A>Status: preliminary; translated from GB/EMBL/DBJ  
A:Molecule type: DNA  
A:Residues: 1-623 <LYN>  
A:Cross-references: EMBL:AL049559; PIDN:CAB40176.1; GSPDB:GNO0068; SPDB:SPCCI450.09C  
A:Experimental source: strain 972h-; cosmid c1450  
C:Genetics:  
A:Gene: SPDB:SPCCI450.09C  
A:Map position: 3  
C:Superfamily: yeast lysophospholipase

Query Match 63.2%; Score 43; DB 2; Length 623;

C:Species: Methanoblobus tindarius  
C:Accession: AF067892  
C:Gene: mcrA  
C:Feature: CDS  
C:Location: 1-1020 bp  
C:Imported: true  
C:Cross-references: GB:AE006641; NID:g13815599; PTDN:AAK42457.1; GSFDB:GN00155

C;Superfamily: precorrin-6Y methylase CblE

Query Match 58.8%; Score 40; DB 2; Length 227;  
Best Local Similarity 54.5%; Pred. No. 24;  
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;  
QY 3 VVGLSPGEQY 13  
: : : : :  
Db 10 IVGVGPDPEY 20

## RESULT 8

I54459  
MHC H-2K1-k - mouse  
C;Species: Mus musculus (house mouse)  
C;Date: 02-Aug-1996 #sequence\_revision 02-Aug-1996 #text\_change 05-Nov-1999  
C;Accession: I54459  
R;Watts, S.; Davis, A.C.; Goodenow, R.S.  
Immunogenetics 29, 355-357, 1989  
A;Title: Sequence analysis of the C3H H-2K1-k gene: Relationship to the H-2 loci.  
A;Reference number: I54459; MUID:89233303; PMID:2714856  
A;Accession: I54459  
A;Status: preliminary; translated from GB/EMBL/DBJ  
A;Molecule type: DNA  
A;Residues: 1-243 <RES>  
A;Cross-references: GB:M27134; NID:g199435; PIDN:AAA39610.1; PID:g387456  
C;Genetics:  
A;Introns: 22/1; 112/1

Query Match 58.8%; Score 40; DB 2; Length 243;  
Best Local Similarity 77.8%; Pred. No. 26;  
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

## QY 4 VGLSPGEQ 12

Db 216 LGLSPGEE 224

## RESULT 9

F75270  
cytochrome P450 - Deinococcus radiodurans (strain R1)  
C;Species: Deinococcus radiodurans  
C;Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000  
C;Accession: F75270  
R;White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;  
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; Ma  
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.  
Science 286, 1571-1577, 1999  
A;Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.  
A;Reference number: A75250; MUID:20036896; PMID:10567266  
A;Accession: F75270  
A;Status: preliminary  
A;Molecule type: DNA  
A;Residues: 1-381 <WHI>  
A;Cross-references: GB:AE002076; GB:AE000513; NID:g6460285; PIDN:AAF12016.1; PID:g646029  
A;Experimental source: strain R1  
C;Genetics:  
A;Gene: DR2473  
A;Map position: 1

Query Match 58.8%; Score 40; DB 2; Length 381;  
Best Local Similarity 100.0%; Pred. No. 41;  
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

## QY 1 PAVVGLSP 8

Db 53 PAVVGLSP 60

## RESULT 10

A75250  
carboxylesterase, type B - Deinococcus radiodurans (strain R1)  
C;Species: Deinococcus radiodurans

C;Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000  
C;Accession: A75250

R;White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;  
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; Ma  
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.  
Science 286, 1571-1577, 1999

A;Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.  
A;Reference number: A75250; MUID:20036896; PMID:10567266  
A;Accession: A75250  
A;Molecule type: DNA

A;Residues: 1-540 <WHI>

A;Cross-references: GB:AE002092; GB:AE000513; NID:g6460455; PIDN:AAF12163.1; PID:g6460404

A;Experimental source: strain R1

C;Genetics:

A;Gene: DR2626

A;Map position: 1

C;Superfamily: cholinesterase; cholinesterase homology

Query Match 58.8%; Score 40; DB 2; Length 540;  
Best Local Similarity 70.0%; Pred. No. 59;  
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

## QY 1 PAVVGLSPGE 10

Db 512 PQVIGLAPGE 521

## RESULT 11

G84511  
hypothetical protein At2g13900 [imported] - Arabidopsis thaliana  
C;Species: Arabidopsis thaliana (mouse-ear cress)  
C;Date: 02-Feb-2001 #sequence\_revision 02-Feb-2001 #text\_change 02-Feb-2001  
C;Accession: G84511  
R;Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.F.; Town, C.D.; Fujii, C.Y.;  
M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon, L.  
Euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, L.  
Nature 402, 761-768, 1999  
A;Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.  
A;Reference number: A84420; MUID:20083487; PMID:10617197  
A;Accession: G84511  
A;Status: preliminary  
A;Molecule type: DNA  
A;Residues: 1-661 <STO>  
A;Cross-references: GB:AE002093; NID:g6598598; PIDN:AAF18650.1; GSPDB:GN00139  
C;Genetics:  
A;Gene: At2g13900  
A;Map position: 2

Query Match 58.8%; Score 40; DB 2; Length 661;  
Best Local Similarity 53.8%; Pred. No. 72;  
Matches 7; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

## QY 1 PAVVGLSPGEQY 13

Db 437 PLTLKIVPGEQY 449

## RESULT 12

T50281  
probable lysophospholipase (EC 3.1.1.5) precursor SPAC977.09c [similarity] - fission yeast  
C;Species: Schizosaccharomyces pombe  
C;Date: 09-Jun-2000 #sequence\_revision 09-Jun-2000 #text\_change 19-Jan-2001  
C;Accession: T50281; F42738  
R;Zimmermann, W.; Wambutt, R.; McDougall, R.C.; Rajandream, M.A.; Barrell, B.G.  
submitted to the EMBL Data Library, January 2000  
A;Reference number: Z25053  
A;Accession: T50281

A;Status: translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-673 <ZIM>

A;Cross-references: EMBL:AL137130; NID:g6742151; PIDN:CAB69631.1; PID:g6742159; GSPDB:GN

A;Experimental source: strain 972h(-); cosmid c977

R;Yoshioka, S.; Kato, K.; Nakai, K.; Okayama, H.; Nojima, H.

us-09-786-214a-13.rpr

Fri May 7 13:19:31 2004

```
DNA Res. 4, 363-369, 1997
A;Title: Identification of open reading frames in Schizosaccharomyces pombe cDNAs.
A;Reference number: Z17323; MUID:96162722; PMID:950191
A;Accession: T42738
A;Status: preliminary; translated from GH/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 427-673 <YOS>
A;Cross-references: EMBL:D89183; NID:gl749573; PIDN:BAA13845.1; PID:gl749574
A;Experimental source: strain PR745
C;Genetics:
A;Gene: SPDB:SPAC977.09c
A;Map position: 1
A;Introns: 651/3
C;Description:
A;Description: catalyzes the hydrolysis of 2-lysophosphatidylcholine to glycerophosphocholine
A;Superfamily: yeast lysophospholipase
C;Keywords: carboxylic ester hydrolase

Query Match 58.8%; Score 40; DB 2; Length 673;
Best Local Similarity 61.5%; Pred. No. 74;
Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 PAVVGLSPGGEQ 13
DB 83 PASEGLNEGEQSY 95

RESULT 13
S21065
Ig kappa chain V region (anti-RH(D)) - human (fragment)
C;Species: Homo sapiens (man)
C;Date: 22-Nov-1993 #sequence_revision 24-May-1996 #text_change 09-May-1997
C;Accession: S21065
R;Dlouha, A.; Lecroisey, A.; Henschen, A.; Rouger, P.; Keil, B.
Protein Seq. Data Anal. 4, 317-318, 1991
A;Title: Subgroup assignment of a human monoclonal anti-Rh(D) antibody.
A;Reference number: S21065; MUID:92253544; PMID:1812483
A;Accession: S21065
A;Molecule type: protein
A;Residues: 1-43 <DLO>
C;Superfamily: immunoglobulin V region; immunoglobulin homology
C;Keywords: heterotetramer; immunoglobulin

Query Match 57.4%; Score 39; DB 2; Length 43;
Best Local Similarity 63.6%; Pred. No. 6.3;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGGEQ 11
DB 8 PATLSLSPGER 18

RESULT 14
S45441
Ig kappa chain V region - human (fragment)
C;Species: Homo sapiens (man)
C;Date: 27-Jan-1995 #sequence_revision 27-Jan-1995 #text_change 21-Jan-2000
C;Accession: S45441
R;Cox, J.P.L.; Tomlinson, I.M.; Winter, G.
Eur. J. Immunol. 24, 827-836, 1994
A;Title: A directory of human germ-line V(kappa) segments reveals a strong bias in their
A;Reference number: S45324; MUID:94200218; PMID:8149953
A;Accession: S45441
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-96 <COX>
A;Cross-references: EMBL:Z27500
C;Superfamily: immunoglobulin V region; immunoglobulin homology
C;Keywords: heterotetramer; immunoglobulin
F;16-91/Domain: immunoglobulin homology <IMM>

Query Match 57.4%; Score 39; DB 2; Length 96;
Best Local Similarity 63.6%; Pred. No. 15;
```

```
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGGEQ 11
DB 8 PATLSLSPGER 18

RESULT 15
S19975
Ig kappa chain V region (M-T408) - mouse (fragment)
C;Species: Mus musculus (house mouse)
C;Date: 06-Feb-1995 #sequence_revision 06-Feb-1995 #text_change 21-Jan-2000
C;Accession: S19975
R;Weissenborn, W.; Riethmuller, G.; Weiss, E.M.; Rieber, E.P.
submitted to the EMBL Data Library, March 1992
A;Description: Structural characterization of CD4 mAb.
A;Reference number: S19963
A;Accession: S19975
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-103 <WEI>
A;Cross-references: EMBL:X65097; NID:g52296; PIDN:CAA46225.1; PID:g52297
C;Superfamily: immunoglobulin V region; immunoglobulin homology
C;Keywords: heterotetramer; immunoglobulin
F;11-85/Domain: immunoglobulin homology <IMM>

Query Match 57.4%; Score 39; DB 2; Length 103;
Best Local Similarity 63.6%; Pred. No. 16;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGGEQ 11
DB 3 PATLSLSPGER 13

Search completed: May 7, 2004, 12:39:07
Job time : 9.48667 secs
```

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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:27:20 ; Search time 9.6 Seconds  
(without alignments)  
150.299 Million cell updates/sec

Title: US-09-786-214A-15

Perfect score: 75

Sequence: 1 AGLPAVVGLSPGEQE 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 segs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 78:\*

1: PIR1.\*

2: PIR2.\*

3: PIR3.\*

4: PIR4.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	50	66.7	381	2 F75270	cytochrome P450 -
2	43	57.3	659	1 GKBPT4	gene 18 protein -
3	43	57.3	659	2 A60885	tail sheath protei
4	43	57.3	821	2 C84304	DNA helicase [impo
5	42	56.0	326	2 T45226	probable N5.N10-me
6	42	56.0	508	2 E70764	probable cobl prot
7	41	54.7	156	2 F87551	conserved hypothet
8	41	54.7	164	2 A87399	ISCC1, transposase
9	41	54.7	177	2 B72580	hypothetical prote
10	41	54.7	214	2 A83416	hypothetical prote
11	40	53.3	130	2 D72783	hypothetical prote
12	40	53.3	243	2 I54459	MHC H-2K1-k - mous
13	40	53.3	358	1 JQ1870	Au1 protein - toma
14	40	53.3	359	2 S58167	fructose-bisphosph
15	40	53.3	359	2 T48396	fructose-bisphosph
16	40	53.3	428	2 AG1304	uracil permease ho
17	40	53.3	428	2 AG1676	uracil permease ho
18	40	53.3	540	2 A75250	carboxylesterase,
19	40	53.3	563	2 T36580	hypothetical prote
20	40	53.3	623	2 T40991	probable lysophosp
21	40	53.3	740	1 B65136	yhgF protein - Esc
22	40	53.3	740	2 A98160	hypothetical prote
23	40	53.3	740	2 H86005	hypothetical prote
24	40	53.3	776	2 AB0938	probable transcrip
25	40	53.3	1340	2 A39808	proteoglycan core
26	40	53.3	1506	2 JC5985	phosphonositide 3
27	40	53.3	2327	2 T42630	aggreccan - bovine
28	39	52.0	43	2 S21065	Ig kappa chain V r
29	39	52.0	96	2 S45441	Ig kappa chain V r

```

30      39      52.0      103      2      S19975      Ig kappa chain V r
31      39      52.0      106      2      P90070      Ig kappa chain V r
32      39      52.0      106      2      PC4282      Ig kappa chain (an
33      39      52.0      107      2      S57444      Ig kappa chain V-J
34      39      52.0      108      2      C30502      Ig kappa chain V r
35      39      52.0      108      2      S33988      Ig kappa chain V r
36      39      52.0      108      2      G44151      Ig kappa chain V r
37      39      52.0      111      2      S23628      Ig kappa chain V r
38      39      52.0      114      2      S54905      Ig kappa chain V r
39      39      52.0      115      1      K3HUVG      Ig kappa chain pre
40      39      52.0      115      1      KVMSL7      Ig kappa chain pre
41      39      52.0      115      2      S11657      Ig kappa chain pre
42      39      52.0      115      2      G72587      hypothetical prote
43      39      52.0      116      2      B25521      Ig kappa chain pre
44      39      52.0      119      2      S41816      Ig kappa chain V r
45      39      52.0      125      2      S40344      Ig kappa chain V-J

```

#### ALIGNMENTS

```

RESULT 1
F75270
Cytochrome P450 - Deinococcus radiodurans (strain R1)
C;Species: Deinococcus radiodurans
C;Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 17-Mar-2000
C;Accession: F75270
R;White, O.; Eissen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;
S.; Smith, H.O.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; M.
Science 286, 1571-1577, 1999
A;Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.
A;Reference number: A75250; MUID:20036896; PMID:10567266
A;Accession: F75270
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-381 <WHI>
A;Cross-references: GB:AE002076; GB:AB000513; NID:g6460285; PIDN:AAF12016.1; PTD:g64602
A;Experimental source: strain R1
C;Genetics:
A;Gene: DR2473
A;Map position: 1

```

```

Query Match      66.7%; Score 50; DB 2; Length 381;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      2 GLPAVVGLSP 11
      |||||
Db      51 GLPAVVGLSP 60

```

```

RESULT 2
GKBPT4
gene 18 protein - phage T4
N;Alternate names: tail sheath protein gp18
C;Species: phage T4
A;Note: host Escherichia coli
C;Date: 30-Sep-1991 #sequence_revision 30-Sep-1991 #text_change 23-Jul-1999
C;Accession: JF0021
R;Aisaka, F.; Nakako, T.; Takahashi, H.; Ishii, S.
J. Virol. 62, 1186-1193, 1988
A;Title: Nucleotide sequence of the tail sheath gene of bacteriophage T4 and amino acid
A;Reference number: JF0021; MUID:88155753; PMID:2964531
A;Accession: JF0021
A;Molecule type: DNA
A;Residues: 1-659 <ARI>
A;Cross-references: GB:M19085; EMBL:M36959; NID:g215949; PIDN:AAA32541.1; PTD:g215950
C;Genetics:
A;Gene: 18
A;Map position: 97.352-99.328
C;Superfamily: phage T4 gene 18 protein

```

Query Match 57.3%; Score 43; DB 1; Length 659;  
 Best Local Similarity 66.7%; Pred. No. 56;  
 Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVGSLPGE 13  
 | : | | | | | | |  
 DB 218 GIPGVVALYPGE 229

RESULT 3  
 A50885  
 tail sheath protein - phage T4  
 C:Species: Halobacterium sp. NRC-1  
 C:Date: 09-Sep-1994 #sequence\_revision 09-Sep-1994 #text\_change 03-May-1996  
 C:Accession: A60885  
 R:Arisaka, F.; Nakako, T.; Kumazaki, T.; Ishii, S.  
 J. Protein Chem. 6, 245-251, 1987  
 A:Title: Primary structure of the tail sheath protein of bacteriophage T4 and its gene.  
 A:Reference number: A60885  
 A:Accession: A60885  
 A:Status: preliminary  
 A:Molecule type: protein  
 A:Residues: 1-659 <ARI>  
 C:Superfamily: phage T4 gene 18 protein

Query Match 57.3%; Score 43; DB 2; Length 659;  
 Best Local Similarity 66.7%; Pred. No. 56;  
 Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVGSLPGE 13  
 | : | | | | | | |  
 DB 218 GIPGVVALYPGE 229

RESULT 4  
 C84304  
 DNA helicase [imported] - Halobacterium sp. NRC-1  
 C:Species: Halobacterium sp. NRC-1  
 C:Date: 02-Feb-2001 #sequence\_revision 02-Feb-2001 #text\_change 02-Feb-2001  
 C:Accession: C84304  
 R:Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.; Leithauer, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jablon, K.H.; Alam, M.; Freitas, T.  
 Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000  
 A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li  
 A:Title: Genome sequence of Halobacterium species NRC-1  
 A:Reference number: A84160; MUID:20504483; PMID:11016950  
 A:Accession: C84304  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-821 <STO>  
 A:Cross-references: GB:AE004437; NID:g10580995; PIDN:AAG19799.1; GSPDB:GN00138  
 C:Genetics:  
 A:Gene: hel

Query Match 57.3%; Score 43; DB 2; Length 821;  
 Best Local Similarity 90.0%; Pred. No. 70;  
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 0;

QY 5 AVVGLSPGEQ 14  
 | : | | | | | | |  
 DB 326 AVVGLSPAQQ 335

RESULT 5  
 T45226  
 Probable N5,N10-methylene-tetrahydromethanopterin reductase (F420-dependent) [imported]  
 C:Species: Methanobolus tindarius  
 C:Date: 21-Jan-2000 #sequence\_revision 21-Jan-2000 #text\_change 21-Jan-2000  
 C:Accession: T45226  
 R:Westenberg, D.J.; Braune, A.; Ruppert, C.; Mueller, V.; Herzberg, C.; Gottschalk, G.;  
 submitted to the EMBL Data Library, September 1998  
 A:Description: The F420H2-dehydrogenase from Methanobolus tindarius: Cloning of the ffd

A:Reference number: Z22947  
 A:Accession: T45226  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-326 <WES>  
 A:Cross-references: EMBL:AJ011519; PIDN:CAB56639.1  
 A:Experimental source: DSM 2278  
 C:Genetics:  
 A:Gene: ffdA

Query Match 56.0%; Score 42; DB 2; Length 326;  
 Best Local Similarity 57.1%; Pred. No. 40;  
 Matches 8; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGLPAVVGSLPGEQ 14  
 | : | : | | | | |  
 DB 84 SGGRAILLGSPGEQ 97

RESULT 6  
 E70764  
 probable cobi protein - Mycobacterium tuberculosis (strain H37RV)  
 C:Species: Mycobacterium tuberculosis  
 C:Date: 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change 22-Oct-1999  
 C:Accession: E70764  
 R:Coile, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, J.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.; Nature 393, 537-544, 1998  
 A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
 A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome  
 A:Reference number: A70500; MUID:98295987; PMID:9634230  
 A:Accession: E70764  
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-508 <COL>  
 A:Cross-references: GB:273966; GB:AL123456; NID:g3261577; PIDN:CAA98214.1; PID:e246996;  
 A:Experimental source: strain H37RV  
 C:Genetics:  
 A:Gene: cobi

Query Match 56.0%; Score 42; DB 2; Length 508;  
 Best Local Similarity 57.1%; Pred. No. 62;  
 Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 GLPAVVGSLPGEQ 15  
 | : | | | | | : |  
 DB 247 GTVAVVGLGPGDSD 260

RESULT 7  
 F87551  
 conserved hypothetical protein CC2439 [imported] - Caulobacter crescentus  
 C:Species: Caulobacter crescentus  
 C:Date: 20-Apr-2001 #sequence\_revision 20-Apr-2001 #text\_change 10-May-2001  
 C:Accession: F87551  
 R:Nierman, W.C.; Feidblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.  
 B.; Laub, M.T.; Deboy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.D.; Haft, D.H.; Kolo  
 n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Shapero, L.; Venter, J.C.; Fraser, C.M.  
 Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001  
 A:Title: Complete Genome Sequence of Caulobacter crescentus.  
 A:Reference number: A87249; MUID:21173698; PMID:11259647  
 A:Accession: F87551  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-156 <STO>  
 A:Cross-references: GB:AE005673; NID:g13423984; PIDN:AAK24410.1; GSPDB:GN00148  
 C:Genetics:  
 A:Gene: CC2439  
 C:Superfamily: Haemophilus influenzae conserved hypothetical protein HI0305

Query Match 54.7%; Score 41; DB 2; Length 156;  
 Best Local Similarity 80.0%; Pred. No. 28;

Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 5 AVVGLSPGEQ 14  
 |||||  
 Db 18 AVVGLDPGEK 27

## RESULT 8

A87399  
 ISCE1, transposase OrfB [imported] - Caulobacter crescentus  
 C:Species: Caulobacter crescentus  
 C>Date: 20-Apr-2001 #sequence\_revision 20-Apr-2001 #text\_change 18-Jul-2001  
 C:Accession: A87399; E87402; E87502; E87570  
 R:Nierman, W.C.; Feidblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.  
 n, J.; Ermolaeva, M.; White, O.; Salzman, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolon  
 Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001  
 A:Title: Complete Genome Sequence of *Caulobacter crescentus*.  
 A:Reference number: A87249; MUID:21173698; PMID:11259647  
 A:Accession: A87399  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-164 <STO>  
 A:Cross-references: GB:AE005673; NID:gl3422533; PIDN:AAK23189.1; GSPDB:GN00148  
 A:Accession: E87402  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-164 <ST2>  
 A:Cross-references: GB:AE005673; NID:gl3422563; PIDN:AAK23217.1; GSPDB:GN00148  
 A:Accession: E87502  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-164 <ST3>  
 A:Cross-references: GB:AE005673; NID:gl3423519; PIDN:AAK24017.1; GSPDB:GN00148  
 A:Accession: E87570  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-164 <ST4>  
 A:Cross-references: GB:AE005673; NID:gl3424163; PIDN:AAK24561.1; GSPDB:GN00148  
 C:Genetics:  
 C:Superfamily: Streptomyces coelicolor probable transposase SC69.35c

Query Match 54.7%; Score 41; DB 2; Length 164;  
 Best Local Similarity 53.8%; Pred. No. 29;  
 Matches 7; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 GLPAVVLSPGEQ 14  
 |||||  
 Db 49 GLPVRLSPGEQ 61

## RESULT 9

B72580  
 hypothetical protein APE1922 - Aeropyrum pernix (strain K1)  
 C:Species: Aeropyrum pernix  
 C>Date: 20-Aug-1999 #sequence\_revision 20-Aug-1999 #text\_change 20-Aug-1999  
 C:Accession: B72580  
 R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takai  
 awa, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K  
 DNA Res. 6, 83-101, 1999  
 A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyr  
 A:Reference number: A72450; MUID:99310339; PMID:10382966  
 A:Accession: B72580  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-177 <KAW>  
 A:Cross-references: DDBJ:AP000062; NID:g5105244; PIDN:BAA80927.1; PID:d1044713; PID:g510  
 A:Experimental source: strain K1  
 C:Genetics:  
 A:Gene: APE1922

Query Match 54.7%; Score 41; DB 2; Length 177;

Best Local Similarity 53.8%; Pred. No. 31;  
 Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 GLPAVVLSPGEQ 14  
 |||||  
 Db 151 GVPVGLSPGEQ 163

## RESULT 10

A83416  
 hypothetical protein PA1825 [imported] - *Pseudomonas aeruginosa* (strain PAO1)  
 C:Species: *Pseudomonas aeruginosa*  
 C>Date: 15-Sep-2000 #sequence\_revision 15-Sep-2000 #text\_change 31-Dec-2000  
 C:Accession: A83416  
 R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.; B  
 adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.K.; Kas, A.; Larbig, K.; Lim  
 ; Lozy, S.; Olson, M.V.  
 Nature 406, 959-964, 2000  
 A:Title: Complete genome sequence of *Pseudomonas aeruginosa* PAO1, an opportunistic path  
 A:Reference number: A82950; MUID:20437337; PMID:10984043  
 A:Accession: A83416  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-214 <STO>  
 A:Cross-references: GB:AE004609; GB:AE004091; NID:g9947810; PIDN:AAG05214.1; GSPDB:GN00  
 A:Experimental source: strain PAO1  
 C:Genetics:  
 A:Gene: PA1825

Query Match 54.7%; Score 41; DB 2; Length 214;  
 Best Local Similarity 63.6%; Pred. No. 38;  
 Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GLPAVVLSPG 12  
 |||||  
 Db 113 GLAALIGLAPG 123

## RESULT 11

D72783  
 hypothetical protein APE0253 - *Aeropyrum pernix* (strain K1)  
 C:Species: *Aeropyrum pernix*  
 C>Date: 20-Aug-1999 #sequence\_revision 20-Aug-1999 #text\_change 09-Jun-2000  
 C:Accession: D72783  
 R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takai  
 awa, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K  
 DNA Res. 6, 83-101, 1999  
 A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyr  
 A:Reference number: A72450; MUID:99310339; PMID:10382966  
 A:Accession: D72783  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-130 <KAW>  
 A:Cross-references: DDBJ:AP000058; NID:g5103388; PIDN:BAA79166.1; PID:d1042942; PID:g51  
 A:Experimental source: strain K1  
 C:Genetics:  
 A:Gene: APE0253  
 C:Superfamily: Aeropyrum pernix hypothetical protein APE0253

Query Match 53.3%; Score 40; DB 2; Length 130;  
 Best Local Similarity 58.3%; Pred. No. 33;  
 Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVLSPGE 13  
 |||||  
 Db 5 GVPVGLSPGE 16

## RESULT 12

I54459  
 MHC H-2K1-k - mouse  
 C:Species: *Mus musculus* (house mouse)  
 C>Date: 02-Aug-1996 #sequence\_revision 02-Aug-1996 #text\_change 05-Nov-1999



C:Accession: I54459  
 R:Watts, S.; Davis, A.C.; Goodenow, R.S.  
 Immunogenetics 29, 355-357, 1989  
 A:Title: Sequence analysis of the C3H H-2K1-k gene: Relationship to the H-2 loci.  
 A:Reference number: I54459; MUID:89233303; PMID:2714856  
 A:Accession: I54459  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-243 <RES>  
 A:Cross-references: GB:M27134; NID:g199435; PIDN:AAA39610.1; PID:g387456  
 C:Genetics:  
 A:Introns: 22/1; 112/1

Query Match 53.3%; Score 40; DB 2; Length 243;  
 Best Local Similarity 77.8%; Pred. No. 62;  
 Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 7 VGLSPGEEQ 15  
 :|||:|:  
 Db 216 LGLSPGEEE 224

RESULT 13  
 JQ1870  
 A11 protein - tomato mottle virus (isolate Florida)  
 C:Species: tomato mottle virus  
 C:Date: 17-Feb-1994 #sequence\_revision 17-Feb-1994 #text\_change 07-May-1999  
 C:Accession: JQ1870  
 R:Abouzid, A.M.; Polston, J.E.; Hiebert, E.  
 J. Gen. Virol. 73, 3225-3229, 1992  
 A:Title: The nucleotide sequence of tomato mottle virus, a new geminivirus isolated from  
 A:Reference number: JQ1869; MUID:93107858; PMID:1469361  
 A:Accession: JQ1870  
 A:Status: translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-358 <ABO>  
 A:Cross-references: GB:L14460  
 C:Genetics:  
 A:Map position: segment A  
 C:Superfamily: tomato golden mosaic virus A11 protein

Query Match 53.3%; Score 40; DB 1; Length 358;  
 Best Local Similarity 58.3%; Pred. No. 92;  
 Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 GLPAVVGLSGPE 13  
 :|||:|:  
 Db 296 GIPAVLCNPGE 307

RESULT 14  
 S58167  
 fructose-bisphosphate aldolase (EC 4.1.2.13) - garden pea  
 A:Alternate names: fructose-1,6-bisphosphate aldolase  
 C:Species: Pisum sativum (garden pea)  
 C:Date: 13-Jan-1996 #sequence\_revision 01-Mar-1996 #text\_change 22-Jun-1999  
 C:Accession: S58167  
 R:Pelzer-Reith, B.; Schnarrenberger, C.  
 submitted to the EMBL Data Library, July 1995  
 A:Description: Characterization of cDNA clones and expression of two cytosolic fructose-  
 A:Reference number: S58167  
 A:Accession: S58167  
 A:Status: preliminary  
 A:Molecule type: mRNA  
 A:Residues: 1-359 <PEL>  
 A:Cross-references: EMBL:X89829; NID:g927504; PIDN:CAA61947.1; PID:g927505  
 C:Superfamily: fructose-bisphosphate aldolase  
 C:Keywords: aldehyde-lyase; carbon-carbon lyase

Query Match 53.3%; Score 40; DB 2; Length 359;  
 Best Local Similarity 60.0%; Pred. No. 92;  
 Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AGLPAVVGLSGPEQ 15  
 :|||:|:  
 Db 258 AAVPAVVGLSGGQSE 272

RESULT 15  
 T48396  
 fructose-bisphosphate aldolase-like protein - Arabidopsis thaliana  
 N:Alternate names: protein F17C15.110  
 C:Species: Arabidopsis thaliana (mouse-ear cress)  
 C:Date: 20-Apr-2000 #sequence\_revision 20-Apr-2000 #text\_change 19-May-2000  
 C:Accession: T48396  
 R:Revan, M.; Pohl, T.; Weizenegger, T.; Bancroft, I.; Mewes, H.W.; Rudd, S.; Lemcke, K.  
 submitted to the Protein Sequence Database, March 2000  
 A:Reference number: Z24492  
 A:Accession: T48396  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-359 <BEV>  
 A:Cross-references: EMBL:AL162506  
 A:Experimental source: cultivar Columbia; BAC clone F17C15  
 C:Genetics:  
 A:Map position: 5  
 A:Introns: 10/1; 86/1  
 A:Note: F17C15.110  
 C:Superfamily: fructose-bisphosphate aldolase

Query Match 53.3%; Score 40; DB 2; Length 359;  
 Best Local Similarity 60.0%; Pred. No. 92;  
 Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AGLPAVVGLSGPEQ 15  
 :|||:|:  
 Db 258 AAVPAVVGLSGGQSE 272

Search completed: May 7, 2004, 12:39:07  
 Job time : 9.76667 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 4.68 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-14  
Perfect score: 65  
Sequence: 1 LPAVVGLSPGGEQ 13

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	63.1	156	1	RVUX_CAUCR
2	41	63.1	326	1	MER_METTI
3	40	61.5	633	1	PLB5_SCHPO
4	39	60.0	115	1	KV3I_HUMAN
5	39	60.0	115	1	KV5I_MOUSE
6	39	60.0	508	1	COBI_MYCTU
7	39	60.0	597	1	NR41_RAT
8	39	60.0	1402	1	N160_MOUSE
9	39	60.0	1636	1	BUD3_YEAST
10	38	58.5	390	1	COBL_MYCTU
11	38	58.5	429	1	RS1_LEULA
12	38	58.5	507	1	CAT4_PICAN
13	38	58.5	516	1	C4AD_DROME
14	38	58.5	699	1	EFG_HAEIN
15	38	58.5	700	1	EFG_PASMU
16	37.5	57.7	827	1	M4K1_MOUSE
17	37	56.9	280	1	PMXA_MOUSE
18	37	56.9	281	1	PMXA_RAT
19	37	56.9	446	1	COBJ_ARCFU
20	37	56.9	446	1	ENOI_MAIZE
21	37	56.9	557	1	PUR6_VIGAC
22	37	56.9	637	1	MUTL_CAUCR
23	37	56.9	658	1	VG18_BPT4
24	37	56.9	813	1	CADM_MOUSE
25	37	56.9	985	1	4ET_HUMAN
26	37	56.9	2269	1	RKSL_SV41
27	36.5	56.2	374	1	RRSG_BOVIN
28	36	55.4	129	1	KV3H_HUMAN
29	36	55.4	211	1	COBL_METJA
30	36	55.4	279	1	TYSY_CAUCR
31	36	55.4	342	1	HOPK_AZOVI
32	36	55.4	348	1	HOPV_AZOVI
33	36	55.4	359	1	ALF2_PEA

34 36 55.4 359 1 ALF\_CICAR  
35 36 55.4 590 1 MUTL\_THETN  
36 36 55.4 598 1 NR41\_CANFA  
37 36 55.4 750 1 ELS\_CHICK  
38 35.5 54.6 833 1 M4K1\_HUMAN  
39 35 53.8 100 1 KV3C\_HUMAN  
40 35 53.8 109 1 KV3B\_HUMAN  
41 35 53.8 109 1 KV3D\_HUMAN  
42 35 53.8 109 1 KV3E\_HUMAN  
43 35 53.8 109 1 KV3G\_HUMAN  
44 35 53.8 129 1 KV3L\_HUMAN  
45 35 53.8 129 1 KV3M\_HUMAN

O65735 cicer ariet  
Q8ra70 thermoanaer  
P51666 canis famil  
Q07916 gallus gall  
Q92918 homo sapien  
P01621 homo sapien  
P01620 homo sapien  
P01622 homo sapien  
P01623 homo sapien  
P04206 homo sapien  
P18135 homo sapien  
P18136 homo sapien

## ALIGNMENTS

RESULT 1  
ID RVUX\_CAUCR STANDARD; PRT; 156 AA.  
AC Q9A5K8;  
DT 10-OCT-2003 (Rel. 42, Created)  
DT 10-OCT-2003 (Rel. 42, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Putative Holliday Junction resolvase (EC 3.1.1.-).  
GN CC2439.  
OS Caulobacter crescentus.  
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;  
OC Caulobacteraceae; Caulobacter.  
OX NCBI\_TaxID=155892;  
RN [1]

SEQUENCE FROM N.A.  
RC STRAIN=ATCC 19089 / CB15;  
RX MEDLINE=21173698; PubMed=11259647;  
RA Nierman W.C., Feldblyum T.V., Laub M.T., Paulsen I.T., Nelson K.E., Eisen J.A., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R., Petočka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B., DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.L., Haft D.H., Kolonay J.F., Smit J., Craven M.B., Khouri H., Shetty J., Berry K., Utterback T., Tran K., Wolf A., Vamathevan J., Ermolaeva M., White O., Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;  
RA "Complete genome sequence of Caulobacter crescentus.";  
RT Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001).  
EL -!- FUNCTION: Could be a nuclease that resolves Holliday junction intermediates in genetic recombination.  
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (Potential).  
CC -!- SIMILARITY: Belongs to the YGF HJR family.  
CC -----

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EMBL; AE005913; AAK24410.1; -;  
DR PIR; F87551; F87551.  
DR TIGR; CC2439; -;  
DR HAMAP; MF\_00651; -; 1.  
DR InterPro; IPR005227; Cons\_hypoth250.  
DR InterPro; IPR006641; YgfC.  
DR Pfam; PF03652; UPF0081; 1.  
DR SMART; SM00732; YGFPC; 1.  
DR TIGRFAMs; TIGR00250; TIGR00250; 1.  
KW Hydrolase; Nuclease; DNA repair; DNA recombination; Complete proteome.  
SQ SEQUENCE 156 AA; 17142 MW; 21F54D8648396141 CRC64;

Query Match 63.1%; Score 41; DB 1; Length 156;  
Best Local Similarity 80.0%; Pred. No. 6.8;  
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGGEQ 12

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Db      18 AVVGLDPGEK 27
|||||
|||||

RESULT 2
MER_METTI
ID MER_METTI STANDARD; PRT; 326 AA.
AC Q9UXP0;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Coenzyme F420-dependent N(S),N(10)-methylentetrahydromethanopterin
DE reductase (EC 1.5.99.11) (Methylene-H(4)MPT reductase).
GN MER OR PFDA.
OS Methanobolus tindarius.
OC Archaea; Euryarchaeota; Methanomicrobia; Methanosarcinales;
OC Methanosarcinaceae; Methanobolus.
OX NCBI_TaxID=2221;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM 2278;
RA Westernberg D.J., Brune A., Ruppert C., Mueller V., Herzberg C.,
RA Gottschalk G., Blaut M.;
RT "The F420H2-dehydrogenase from Methanobolus tindarius: cloning of the
RT ffd operon and expression of the genes in Escherichia coli.";
RL FEMS Microbiol. Lett. 170:389-398(1999).
CC -!- FUNCTION: Catalyzes the reversible reduction of methylene-H(4)MPT
CC to methyl-H(4)MPT (By similarity).
CC -!- CATALYTIC ACTIVITY: N(5),N(10)-methylentetrahydromethanopterin +
CC reduced coenzyme F420 = 5-methyl-5,6,7,8-tetrahydromethanopterin +
CC coenzyme F420.
CC -!- PATHWAY: Methanogenesis from carbon dioxide; fifth step.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the mer family.
CC
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CC
CC EMBL; AJ011519; CAB56639.1; -.
CC F1R; T45226; T45226.
CC HAMAP; MF 01091; -.
CC InterPro; IPR002103; Bac_luciferase.
CC Pfam; PF00296; bac_luciferase; 1.
KW Methanogenesis; One-carbon metabolism; Oxidoreductase.
SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;

Query Match 63.1%; Score 41; DB 1; Length 326;
Best Local Similarity 70.0%; Pred. No. 14;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGEQ 12
|::| |
|::| |

Db 88 AILGLGPEQ 97

RESULT 3
PLB5 SCHPO STANDARD; PRT; 633 AA.
ID PLB5 SCHPO
AC Q9Y7N6;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Putative lysophospholipase C1450.09c precursor (EC 3.1.1.5)
DE (Phospholipase B).
DE SPOC1450.09c.
GN Schizosaccharomyces pombe (Fission yeast).
OS Schizosaccharomycetes; Ascomycota; Schizosaccharomycetes;
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;

OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21849401; PubMed=11859360;
RA Wood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgouros J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Fellwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volckaert G., Aert R., Robben J., Grymonprez B.,
RA Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Moestl D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gallard C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerretti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe.";
RT Nature 415:871-880(2002).
RL Nature 415:871-880(2002).
CC -!- FUNCTION: Catalyzes the release of fatty acids from
CC lysophospholipids (By similarity).
CC -!- CATALYTIC ACTIVITY: 2-lysophosphatidylcholine + H(2)O =
CC glycerophosphocholine + a fatty acid anion.
CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
CC -!- SIMILARITY: Belongs to the lysophospholipase family.
CC
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CC
CC EMBL; AL049559; CAB40176.2; -.
CC GenDB SPombe; SPOC1450.09c; -.
CC InterPro; IPR002642; PLAC.
CC Pfam; PF01735; PLA2_B; 1.
CC SMART; SM00022; PLAC; 1.
CC Hypothetical protein; Lipid degradation; Hydrolase; Glycoprotein;
CC Signal.
CC
CC SIGNAL 1 19 POTENTIAL.
CC CHAIN 20 633 PUTATIVE LYSOPHOSPHOLIPASE C1450.09c.
CC CARBOHYD 118 118 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 153 153 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 187 187 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 232 232 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 256 256 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 264 264 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 293 293 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 331 331 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 360 360 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 367 367 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 400 400 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 403 403 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 474 474 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 508 508 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 513 513 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 537 537 N-LINKED (GLCNAC. .) (POTENTIAL).

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FT CARBOHYD 564 586 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 586 586 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 603 603 N-LINKED (GLCNAC. . .) (POTENTIAL).  
SQ SEQUENCE 633 AA; 68292 MW; 49871B2955893D19 CRC64;

Query Match 61.5%; Score 40; DB 1; Length 633;  
Best Local Similarity 75.0%; Pred. No. 39;  
Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 13  
||| ||| ||| |||  
DB 76 PASDGLSTGEQ 87

RESULT 4  
KV31\_HUMAN STANDARD; PRT; 115 AA.  
AC P04433;  
DT 13-AUG-1987 (Rel. 05, Created)  
DT 13-AUG-1987 (Rel. 05, Last sequence update)  
DT 15-JUL-1999 (Rel. 38, Last annotation update)  
DE IG kappa chain V-III region VG precursor (Fragment).  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
RP SEQUENCE FROM N.A. PubMed=6440122;  
RX MEDLINE=85087932; PubMed=6440122;  
RA Pech M., Zachau H.G.;  
RT "Immunoglobulin genes of different subgroups are interdigitated within the VK locus.";  
RL Nucleic Acids Res. 12:9229-9236(1984).  
CC

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EMBL; X01668; -; NOT ANNOTATED\_CDS.  
PIR; A01900; K3HUVG.  
HSP; P80362; 1WTL.

GO; GO:0005576; C:extracellular; NAS.  
GO; GO:0003823; F:antigen binding; NAS.  
GO; GO:0006955; F:immune response; NAS.  
InterPro; IPR007110; IG-like.  
InterPro; IPR003596; IG\_v.  
Pfam; PF00047; IG; 1.  
SMART; SM00406; IGv; 1.  
PROSITE; PS00835; IG LIKE; 1.  
Immunoglobulin V region; Signal.

FT SIGNAL 1 20  
FT CHAIN 21 >115 IG KAPPA CHAIN V-III REGION VG.  
FT DOMAIN 21 43 FRAMEWORK-1.  
FT DOMAIN 44 54 COMPLEMENTARITY-DETERMINING-1.  
FT DOMAIN 55 69 FRAMEWORK-2.  
FT DOMAIN 70 76 COMPLEMENTARITY-DETERMINING-2.  
FT DOMAIN 77 108 FRAMEWORK-3.  
FT DOMAIN 109 115 COMPLEMENTARITY-DETERMINING-3.  
FT DISULFID 43 108 BY SIMILARITY.  
FT NON TER 115 115  
SQ SEQUENCE 115 AA; 12575 MW; 2DE47CDA3A17D555 CRC64;

Query Match 60.0%; Score 39; DB 1; Length 115;  
Best Local Similarity 63.6%; Pred. No. 11;  
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
||| : |||||  
DB 28 PATLSVSPGER 38

RESULT 6  
COBI\_MYCTU STANDARD; PRT; 508 AA.  
ID COBI\_MYCTU  
AC Q10677;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Cobalamin biosynthesis protein COBIJ [includes: Precorrin-2 C20-methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2 methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.1.-)].  
GN COBIJ OR COBI OR RV2066 OR MT2126 OR MTCY49.05 OR MB2092.  
OS Mycobacterium tuberculosis, and  
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
OC Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.  
OX NCBI\_TaxID=1773, 1765;  
RN [1]  
RP SEQUENCE FROM N.A.

RESULT 5  
KV51\_MOUSE STANDARD; PRT; 115 AA.  
AC P01642;  
DT 21-JUL-1986 (Rel. 01, Created)  
DT 21-JUL-1986 (Rel. 01, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE IG kappa chain V-V region L7 precursor (Fragment).  
OS Mus musculus (Mouse).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
OX NCBI\_TaxID=10090;  
RN [1]  
RP SEQUENCE FROM N.A. PubMed=6264318;  
RX MEDLINE=81220975; PubMed=6264318;  
RA Pech M., Hochtl J., Schnell H., Zachau H.G.;  
RT "Differences between germ-line and rearranged immunoglobulin V kappa coding sequences suggest a localized mutation mechanism.";  
RL Nature 291:668-670(1981).  
CC -!- MISCELLANEOUS: THERE APPEAR TO BE TWO POSSIBLE SPICE JUNCTIONS AT THE 3' END OF THE INTRON. THE ALTERNATE WOULD CODE FOR A PROTEIN LACKING RESIDUES 17-19.

PIR; A01925; KVMSL7.  
PDB; 1J10; 18-FEB-03.  
PDB; 1J1P; 18-FEB-03.  
PDB; 1J1X; 18-FEB-03.  
InterPro; IPR007110; IG-like.  
InterPro; IPR003596; IG\_v.  
Pfam; PF00047; IG; 1.  
SMART; SM00406; IGv; 1.  
PROSITE; PS00835; IG LIKE; 1.  
Immunoglobulin V region; Signal; 3D-structure.

FT SIGNAL 1 20  
FT CHAIN 21 >115 IG KAPPA CHAIN V-V REGION L7.  
FT DOMAIN 21 43 FRAMEWORK-1.  
FT DOMAIN 44 54 COMPLEMENTARITY-DETERMINING-1.  
FT DOMAIN 55 69 FRAMEWORK-2.  
FT DOMAIN 70 76 COMPLEMENTARITY-DETERMINING-2.  
FT DOMAIN 77 108 FRAMEWORK-3.  
FT DOMAIN 109 >115 COMPLEMENTARITY-DETERMINING-3.  
FT DISULFID 43 108 BY SIMILARITY.  
FT NON TER 115 115  
SQ SEQUENCE 115 AA; 12615 MW; C17BEC758C577E00 CRC64;

Query Match 60.0%; Score 39; DB 1; Length 115;  
Best Local Similarity 54.5%; Pred. No. 11;  
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
||| : |||||  
DB 28 PATLSVSPGER 38

RESULT 6  
COBI\_MYCTU STANDARD; PRT; 508 AA.  
ID COBI\_MYCTU  
AC Q10677;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Cobalamin biosynthesis protein COBIJ [includes: Precorrin-2 C20-methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2 methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.1.-)].  
GN COBIJ OR COBI OR RV2066 OR MT2126 OR MTCY49.05 OR MB2092.  
OS Mycobacterium tuberculosis, and  
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
OC Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.  
OX NCBI\_TaxID=1773, 1765;  
RN [1]  
RP SEQUENCE FROM N.A.

RC SPECIES=M.tuberculosis; STRAIN=H37Rv;  
RX MEDLINE=98235987; PubMed=9634230;  
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,  
Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,  
Badcock K., Badham D., Brown D., Chillingworth T., Connor R.,  
Davies R., Devlin K., Fellwell T., Gentles S., Hamlin N., Holroyd S.,  
Hornshy T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,  
Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,  
Rutter S., Seeger K., Skelton S., Squares S., Squares R.,  
Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;  
RA "Deciphering the biology of Mycobacterium tuberculosis from the  
RT complete genome sequence.";  
RL Nature 393:537-544 (1998).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC SPECIES=M.tuberculosis; STRAIN=CDC 1551 / Oshkosh;  
RX MEDLINE=2226494; PubMed=12218036;  
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,  
Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,  
Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,  
Delcher A., Uterback T., Weidman J., Khouri H., Gill J., Mikula A.,  
Biswal W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;  
RA "Whole-genome comparison of Mycobacterium tuberculosis clinical and  
RT laboratory strains.";  
RL J. Bacteriol. 184:5479-5490 (2002).  
RN [3]  
RP SEQUENCE FROM N.A.  
RC SPECIES=M.bovis; STRAIN=AF2122/97;  
RX MEDLINE=22709107; PubMed=12788972;  
RA Garnier T., Eiglmeier K., Camus J.-C., Medina N., Mansoor H.,  
Pryor M., Duthoy S., Grondin S., Lacroix C., Monsemp C., Simon S.,  
Harris B., Atkin R., Doggett J., Mayes R., Keating L., Wheeler P.R.,  
Parkhill J., Barrell B.G., Cole S.T., Gordon S.V., Hewinson R.G.;  
RA "The complete genome sequence of Mycobacterium bovis.";  
RL Proc. Natl. Acad. Sci. U.S.A. 100:7877-7882 (2003).  
CC -!- FUNCTION: METHYLATES PRECORRIN-2 AT THE C-20 POSITION TO PRODUCE  
CC PRECORRIN-3A (BY SIMILARITY).  
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + precorrin-2 = S-  
CC adenosyl-L-homocysteine + precorrin-3A.  
CC -!- PATHWAY: Cobalam biosynthesis.  
CC -!- SIMILARITY: BELONGS TO A FAMILY THAT GROUPS SUMT, CYSG, CBIF/COBM  
CC AND CBIL/CBIL.  
CC  
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CC -----  
DR EMBL; Z73966; CAA98214.1; -;  
DR EMBL; AE007063; AAK46406.1; -;  
DR EMBL; EX248341; CAD96945.1; -;  
DR FIR; E70764; E70764.  
DR TIGR; MT2126; -;  
DR TubercuList; RV2066; -;  
DR InterPro; IPR006364; Cobi CbiL.  
DR InterPro; IPR006363; CobiJ.  
DR InterPro; IPR000878; Cor/por Metransf.  
DR InterPro; IPR003043; Uropor Metransf.  
DR Pfam; PF00590; TP methylase; 2.  
DR TIGRFAMs; TIGR01467; cobi cbiL; 1.  
DR TIGRFAMs; TIGR01466; cobi cbiH; 1.  
DR PROSITE; PS00839; SUMT\_1; 1.  
DR PROSITE; PS00840; SUMT\_2; 1.  
DR Cobalamin biosynthesis; Porphyryrin biosynthesis; Transfexase;  
KW Methyltransferase; Multifunctional enzyme; Complete proteome;  
FT DOMAIN 1 243 PRECORRIN-2 C20-METHYLTRANSFERASE.  
FT DOMAIN 244 508 PRECORRIN-3 METHYLASE.  
SQ SEQUENCE 508 AA; 53910 MW; 95AC066F022C4DC1 CRC64;

Query Match

60.0%; Score 39; DB 1; Length 508;

Best Local Similarity 63.6%; Pred. No. 46;  
Matches 7; Conservative 2; Mismatches 0; Gaps 0;  
QY 3 AVUGLSPGEQE 13  
Db 250 AVUGLSPGEQSD 260  
  
RESULT 7  
NR41 RAT  
ID NR41 RAT STANDARD; PRT; 597 AA.  
AC P22829;  
DT 01-AUG-1991 (Rel. 19, Created)  
DT 01-OCT-1996 (Rel. 34, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Orphan nuclear receptor HMR (Nerve growth factor induced protein I-B)  
DE (NGFI-B) (NUR77).  
GN NR4A1 OR HMR OR NGFIB.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
OX NCBI\_TaxID=10116;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=90166506; PubMed=3272167;  
RA Milbrandt J.;  
RT "Nerve growth factor induces a gene homologous to the glucocorticoid  
RT receptor gene.";  
RL Neuron 1:183-188 (1988).  
RN [2]  
RP CHARACTERIZATION.  
RX MEDLINE=93361012; PubMed=8395013;  
RA Wilson T.E., Fahrner T.J., Milbrandt J.;  
RT "The orphan receptors NGFI-B and steroidogenic factor 1 establish  
RT monomer binding as a third paradigm of nuclear receptor-DNA  
RT interaction.";  
RL Mol. Cell. Biol. 13:5794-5804 (1993).  
RN [3]  
RP DNA BINDING MOTIFS.  
RX MEDLINE=92229411; PubMed=1314418;  
RA Wilson T.E., Paulsen R.E., Padgett K.A., Milbrandt J.;  
RT "Participation of non-zinc finger residues in DNA binding by two  
RT nuclear orphan receptors.";  
RL Science 256:107-110 (1992).  
RN [4]  
RP PHOSPHORYLATION.  
RX MEDLINE=94043340; PubMed=8227042;  
RA Hirata Y., Kuchiki K., Chen H.-C., Milbrandt J., Guroff G.;  
RT "The phosphorylation and DNA binding of the DNA-binding domain of the  
RT orphan nuclear receptor NGFI-B.";  
RL J. Biol. Chem. 268:24808-24812 (1993).  
RN [5]  
RX X-RAY CRYSTALLOGRAPHY (2.7 ANGSTROMS) OF 264-351 IN COMPLEX WITH NBRE,  
RP AND METAL-BINDING.  
RX MEDLINE=99260743; PubMed=10331876;  
RA Meinke G., Sigler P.B.;  
RT "DNA-binding mechanism of the monomeric orphan nuclear receptor  
RT NGFI-B.";  
RL Nat. Struct. Biol. 6:471-477 (1999).  
CC -!- FUNCTION: Probable nuclear receptor. May act concomitantly with  
CC NURR1 in regulating the expression of delayed-early genes during  
CC liver regeneration. Binds the NGFI-B response element (NBRE) 5'-  
CC AAAAGGTCA-3'.  
CC -!- SUBUNIT: Binds DNA as a monomer.  
CC -!- SUBCELLULAR LOCATION: Nuclear.  
CC -!- TISSUE SPECIFICITY: Expressed in lung, brain and superior  
CC cervical ganglia. High levels are seen in the adrenal tissue.  
CC -!- INDUCTION: By nerve growth factor and during liver regeneration.  
CC -!- PM: Phosphorylation of Ser-350 results in decrease in NBRE  
CC binding while phosphorylation of Ser-340 has little effect on it.  
CC -!- SIMILARITY: Belongs to the nuclear hormone receptor family. NR4  
CC subfamily.  
CC -----

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CC -----  
 CC EMBL; U17254; AAA56770.1; ALT\_INIT.  
 CC PDB; 1CIT; 26-JUN-00.  
 CC TRANSFAC; T00619; Hormone\_rec\_lig.  
 CC InterPro; IPR000536; Hormone\_rec\_lig.  
 CC InterPro; IPR001723; Strnrm\_receptor.  
 CC InterPro; IPR008946; Strnrm\_receptor.  
 CC InterPro; IPR001628; Znf C4steroid.  
 CC Pfam; PF00104; hormone\_rec; 1.  
 CC Pfam; PF00105; zf-C4; 1.  
 CC PRINTS; PR000398; STRDHOMER.  
 CC PRINTS; PR00047; STROIDFINGER.  
 CC ProDom; PD000035; Znf C4steroid; 1.  
 CC SMART; SM00430; HOL1; 1.  
 CC SMART; SM00399; Znf C4; 1.  
 CC PROSITE; PS00031; NUCLEAR\_RECEPTOR; 1.  
 CC Receptor; Transcription regulation; DNA-binding; Nuclear protein;  
 CC Zinc-finger; Phosphorylation; 3D-structure.  
 CC DNA BIND 266 331 NUCLEAR RECEPTOR-TYPE  
 CC FT ZN\_FING 266 286 C4-TYPE.  
 CC FT ZN\_FING 302 326 C4-TYPE.  
 CC FT DOMAIN 408 458 LIGAND-BINDING (POTENTIAL).  
 CC FT DOMAIN 80 91 POLY-SER.  
 CC FT DOMAIN 182 186 POLY-PRO.  
 CC FT DOMAIN 592 595 PHOSPHORYLATION (BY PKA).  
 CC FT MOD RES 340 340 PHOSPHORYLATION (BY PKA).  
 CC FT MUTAGEN 340 340 S->A: LOSS OF PHOSPHORYLATION.  
 CC FT MUTAGEN 350 350 S->A: LOSS OF PHOSPHORYLATION.  
 CC FT MUTAGEN 345 345 R->K: DECREASED NERE BINDING.  
 CC FT MUTAGEN 348 348 L->V: ALMOST COMPLETE LOSS OF NERE  
 CC BINDING.  
 CC SQ SEQUENCE 597 AA; 64281 MW; 9CFA987112337E53 CRC64;

Query Match 60.0%; Score 39; DB 1; Length 597;  
 Best Local Similarity 46.2%; Pred. No. 54;  
 Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPVVGLSPGEQ 13  
 Db 423 IPGFTELSFGDQ 435

RESULT 8  
 ID N160 MOUSE STANDARD; PRT; 1402 AA.  
 AC Q9Z0W3; Q9CZD9;  
 DT 16-OCT-2001 (Rel. 40, Created)  
 DT 16-OCT-2001 (Rel. 40, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Nuclear pore complex protein Nup160 (Nucleoporin Nup160) (160 kDa  
 DE nucleoporin) (Gene trap locus 1-13) (GTL-13).  
 GN NUP160 OR GTL1-13 OR KIAA0197.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_TaxID=10090;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-129/SVJ;  
 RA Van de Putte T., Cozijnsen M., Dewulf N., Tylzanowski P., Lonnoy O.,  
 RA Huylebroeck D.;  
 RT "Mus musculus mRNA for gtl-13 (gene trap locus-13), similar to human  
 RT KIAA0197 gene (D83781), complete cds";  
 RL Submitted (JUN-1999) to the EMBL/GenBank/DBSJ databases.  
 RN [2]

RP SEQUENCE OF 1151-1402 FROM N.A.  
 RC STRAIN=CS7BL/6J; TISSUE=Embryo;  
 RX MEDLINE=21085660; PubMed=11217851;  
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,  
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,  
 RA Aizawa K., Iwata M., Nishi K., Kiyosawa H., Kondo S., Yamana K.I.,  
 RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.,  
 RA Kadota K., Matsuda H.A., Aeshburner M., Batalov S., Casavant T.,  
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,  
 RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,  
 RA Schriml L.M., Staubli F., Suzuki R., Tonita M., Wagner L., Washio T.,  
 RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,  
 RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,  
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,  
 RA Gustingich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,  
 RA Lyons P., Marchionni L., Mashima J., Marzarelli J., Mombaerts P.,  
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,  
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,  
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,  
 RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.,  
 RA Hayashizaki Y.;  
 RT "Functional annotation of a full-length mouse cDNA collection.";  
 RL Nature 403:685-690(2001).  
 RN [3]  
 RP IDENTIFICATION, SUBUNIT, AND SUBCELLULAR LOCATION.  
 RX MEDLINE=21448620; PubMed=11564755;  
 RA Belgareh N., Rabut G., Bai S.W., van Overbeek M., Beaudouin J.,  
 RA Daigle N., Zatssepina O.V., Pasteau F., Labas V., Fromont-Racine M.,  
 RA Ellenberg J., Doye V.;  
 RT "An evolutionarily conserved NPC subcomplex, which redistributes in  
 RT part to kinetochores in mammalian cells.";  
 RJ J. Cell Biol. 154:1147-1160(2001).  
 RN [4]  
 RP IDENTIFICATION, FUNCTION, SUBUNIT, AND SUBCELLULAR LOCATION.  
 RX MEDLINE=21541555; PubMed=11684705;  
 RA Vasu S., Shah S., Orjalo A., Park M., Fischer W.H., Forbes D.J.;  
 RT "Novel vertebrate nucleoporins Nup133 and Nup160 play a role in mRNA  
 RT export.";  
 RJ J. Cell Biol. 155:339-354(2001).  
 CC -!- FUNCTION: Involved in poly(A)+ RNA transport.  
 CC -!- SUBUNIT: Forms part of the Nup160 subcomplex in the nuclear pore  
 CC which is composed of Nup160, Nup133, Nup107 and Nup96. This  
 CC complex plays a role in RNA export and in tethering Nup98 and  
 CC Nup153 to the nucleus.  
 CC -!- SUBCELLULAR LOCATION: Nuclear pore complex.  
 CC -!- CAUTION: Ref.2 sequence differs from that shown due to a  
 CC frameshift in position 1157 and a stop codon in position 1396.  
 CC -----  
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CC -----  
 CC EMBL; AF104415; AADI7922.2; -  
 CC EMBL; AK012715; BAB28429.1; ALT\_FRAME.  
 CC MGD; MGI:1926227; Nup160.  
 CC GO; GO:0005643; C:nuclear pore; IDA.  
 CC GO; GO:0005487; F:nucleocytoplasmic transporter activity; IDA.  
 CC GO; GO:0006406; P:mRNA-nucleus export; IDA.  
 CC NUCLEAR protein; Transport.  
 CC CONFLICT 1156 1156 A -> T (IN REF. 2).  
 CC CONFLICT 1314 1314 E -> G (IN REF. 2).  
 CC CONFLICT 1368 1368 N -> D (IN REF. 2).  
 CC CONFLICT 1402 1402 AA; 158230 MW; 3BF5D9F057D28772 CRC64;  
 CC SEQUENCE 1402 AA; 158230 MW; 3BF5D9F057D28772 CRC64;

Query Match 60.0%; Score 39; DB 1; Length 1402;  
 Best Local Similarity 70.0%; Pred. No. 1.2e+02;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 LPVVGLSPG 10



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CC      METHYLASES INVOLVED IN COBALAMIN BIOSYNTHESIS.
CC      -----
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CC      between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC      -----
CC      EMBL; Z73966; CAA38225.1; -.
CC      EMBL; AEO07063; AAK46412.1; -.
CC      PIR; C70765; C70765.
CC      TIGR; MT2132; -.
CC      Tuberculin; Rv2072c; -.
CC      InterPro; IPR006365; COBL.
CC      InterPro; IPR000878; Cor/por Metransf.
CC      InterPro; IPR000051; SAM bind.
CC      Pfam; PF00590; TP methylase; 1.
CC      TIGRPFAMs; TIGR01468; COBL cbiET; 1.
CC      Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW      Methyltransferase; Complete proteome.
FT      CONFLICT 205 205 L -> P (IN REF. 2).
FT      CONFLICT 327 327 D -> H (IN REF. 2).
SQ      SEQUENCE 390 AA; 41854 MW; FB42EPF7562F21F3 CRC64;

Query Match      58.5%; Score 38; DB 1; Length 390;
Best Local Similarity 88.9%; Pred. No. 53;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      1 LPAVUGLSP 9
DB      55 LPAVUGLSP 63

RESULT 11
RS1_LEULA
ID      RS1_LEULA STANDARD; PRT; 429 AA.
AC      P50889; P71450;
DT      01-OCT-1996 (Rel. 34, Created)
DT      15-JUL-1999 (Rel. 38, Last sequence update)
DT      16-OCT-2001 (Rel. 40, Last annotation update)
DE      40S ribosomal protein S1.
GN      RPS1
OS      Leuconostoc lactis.
OC      Bacteria; Firmicutes; Lactobacillales; Leuconostoc.
OX      NCBI_TaxID=1246;
RN      [1]
RP      SEQUENCE FROM N.A.
RX      MEDLINE=97186703; PubMed=9034319;
RA      Yamit-Hezi A., Levy Z., Neuman S., Nudel U.;
RT      "A Leuconostoc lactis protein with homology to ribosomal protein S1
RT      shares common epitopes and common DNA binding properties with a
RT      mammalian DNA binding nuclear factor.";
RL      Gene 185:99-103(1997).
RL      [2]
RP      SEQUENCE OF 24-429 FROM N.A.
RX      MEDLINE=95237615; PubMed=7721096;
RA      Eklund E.A., Lee S.W., Skalknik D.G.;
RT      "Cloning of a cDNA encoding a human DNA-binding protein similar to
RT      ribosomal protein S1.";
RL      Gene 155:231-235(1995).
RL      [3]
RP      SEQUENCE OF 78-429 FROM N.A.
RX      MEDLINE=96164600; PubMed=8568274;
RA      Tsuzaka K., Leu A.K., Frank M.B., Movafagh B.F., Koscec M.,
RA      Winkler T.H., Kalden J.R., Reichlin M.;
RT      "Lupus autoantibodies to double-stranded DNA cross-react with
RT      ribosomal protein S1.";
RL      J. Immunol. 156:1668-1675(1996).
CC      -!- FUNCTION: EXHIBITS PREFERENTIAL BINDING TO SINGLE-STRANDED AND
CC      DOUBLE-STRANDED DNA AND A LOW BINDING AFFINITY FOR RNA.
CC      -!- SIMILARITY: Belongs to the S1P family of ribosomal proteins.

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CC      -!- SIMILARITY: Contains 4 S1 motif domains.
CC      -!- CAUTION: WAS ORIGINALLY (REF.2 AND REF.3) THOUGHT TO ORIGINATE
CC      FROM HUMAN BUT IS MOST PROBABLY THE RESULT OF A CDNA LIBRARY
CC      CONTAMINATION BY L.LACTIS.
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CC      or send an email to license@isb-sib.ch).
CC      -----
CC      EMBL; U24086; AAB08978.1; -.
CC      EMBL; U05589; AAA77669.1; -.
CC      EMBL; U27517; AAA97575.1; -.
CC      HSSP; P05055; ISRO.
CC      InterPro; IPR008994; Nucleic acid OB.
CC      InterPro; IPR000110; Ribosomal_S1.
CC      Pfam; PF00575; S1; 4.
CC      PRINTS; PR00681; RIBOSOMALS1.
CC      SMART; SM00316; S1; 4.
CC      PROSITE; PS0126; S1; 4.
KW      Ribosomal protein; Repeat; RNA-binding.
FT      DOMAIN 55 128 S1 MOTIF 1.
FT      DOMAIN 144 211 S1 MOTIF 2.
FT      DOMAIN 231 299 S1 MOTIF 3.
FT      DOMAIN 316 385 S1 MOTIF 4.
FT      CONFLICT 24 24 S -> G (IN REF. 2).
FT      CONFLICT 122 122 A -> S (IN REF. 3).
FT      CONFLICT 217 217 L -> R (IN REF. 2 AND 3).
SQ      SEQUENCE 429 AA; 46386 MW; 92AC82605F39DDFC CRC64;

Query Match      58.5%; Score 38; DB 1; Length 429;
Best Local Similarity 80.0%; Pred. No. 58;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY      3 AVVGLSPGEQ 12
DB      71 AVVGLSTGEE 80

RESULT 12
CATA_PICAN
ID      CATA_PICAN STANDARD; PRT; 507 AA.
AC      P30263;
DT      01-APR-1993 (Rel. 25, Created)
DT      01-APR-1993 (Rel. 25, Last sequence update)
DT      10-OCT-2003 (Rel. 42, Last annotation update)
DE      Peroxisomal catalase (BC 1.11.1.6).
GN      PXP9 OR PXP-9.
OS      Pichia angusta (Yeast) (Hansenula polymorpha).
OC      Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC      Saccharomycetales; Saccharomycetaceae; Pichia.
OX      NCBI_TaxID=4905;
RN      [1]
RP      SEQUENCE FROM N.A.
RX      STRAIN=ATCC 34438;
RX      MEDLINE=92299073; PubMed=1607006;
RA      Didion T., Roggenkamp R.O.;
RT      "Targeting signal of the peroxisomal catalase in the methyiotrophic
RT      yeast Hansenula polymorpha.";
RL      FEBS Lett. 303:113-116(1992).
CC      -!- FUNCTION: Occurs in almost all aerobically respiring organisms and
CC      serves to protect cells from the toxic effects of hydrogen
CC      peroxide.
CC      -!- CATALYTIC ACTIVITY: 2 H(2)O(2) = O(2) + 2 H(2)O.
CC      -!- COFACTOR: Heme group.
CC      -!- SUBUNIT: Homotetramer.
CC      -!- SUBCELLULAR LOCATION: Peroxisomal.
CC      -!- SIMILARITY: Belongs to the catalase family.

```



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EMBL; X56501; CAA39856.1; --  
 DR PIR; S23422; S23422.  
 DR HSSP; F15202; IA4E.  
 DR InterPro; IPR002226; Catalase.  
 DR Pfam; PF00199; catalase; 1.  
 DR PRINTS; PR00067; CATALASE.  
 DR ProDom; PD000510; Catalase; 1.  
 DR PROSITE; PS00437; CATALASE\_1; 1.  
 DR PROSITE; PS00438; CATALASE\_2; 1.  
 DR Oxidoreductase; Peroxidase; Iron; Heme; Hydrogen peroxide;  
 KW Peroxisome.  
 FT ACT SITE 65 BY SIMILARITY.  
 FT ACT SITE 138 138 BY SIMILARITY.  
 FT METAL 348 348 IRON (HEME AXIAL LIGAND) (BY SIMILARITY).  
 FT SITE 505 507 MICROBODY TARGETING SIGNAL (POTENTIAL).  
 SQ SEQUENCE 507 AA; 57849 MW; 35366DDA49539CC3 CRC64;

Query Match 58.5%; Score 38; DB 1; Length 507;  
 Best Local Similarity 70.0%; Pred. No. 68;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
 QY 4 VVGLSPGEQE 13  
 DB 445 VLGRTFGEQE 454  
 |:|:|||||

## RESULT 13

C4AD DROME STANDARD; PRT; 516 AA.  
 AC Q9VAT3;  
 DT 16-OCT-2001 (Rel. 40, Created)  
 DT 16-OCT-2001 (Rel. 40, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Probable cytochrome P450 4adi (EC 1.14.-.-) (CYP14AD1).  
 GN CYP4AD1 OR CG2110.  
 OS Drosophila melanogaster (Fruit fly).  
 OC Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;  
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
 OC Ephydroidea; Drosophilidae; Drosophila.  
 OX NCBI\_TaxID=7227;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=Berkeley;  
 RX MEDLINE=20196006; PubMed=10731132;  
 RA Adams M.D., Celnik S.E., Holt R.A., Evans C.A., Gocayne J.D.,  
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,  
 RA George K.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,  
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,  
 RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,  
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,  
 RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,  
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,  
 RA Beeson D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,  
 RA Burtis K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,  
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,  
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,  
 RA Dodson K., Doup I.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,  
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,  
 RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,  
 RA Glöck A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,  
 RA Harris N.L., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,  
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,  
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,  
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,

LA Lasko P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.,  
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,  
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,  
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,  
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,  
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,  
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,  
 RA Shue B.C., Siden-Kianos I., Simpson M., Skupski M.P., Smith T.,  
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,  
 RA Svirekas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,  
 RA Wang Z.-Y., Wassarman D.A., Weinstock G.M., Weissbach J.,  
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,  
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,  
 RA Zhong X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,  
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;  
 RT "The genome sequence of Drosophila melanogaster.";  
 RL Science 287:2185-2195(2000).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=Berkeley; TISSUE=Embryo;  
 RX MEDLINE=22426066; PubMed=12537569;  
 RA Stapleton M., Carlson J.W., Brokstein P., Yu C., Champe M.,  
 RA George R.A., Guarin H., Kronmiller B., Pacleb J.M., Park S., Wan K.H.,  
 RA Rubin G.M., Celnik S.E.;  
 RT "A Drosophila full-length cDNA resource.";  
 RL Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8(2002).  
 CC -!- FUNCTION: May be involved in the metabolism of insect hormones and  
 CC in the breakdown of synthetic insecticides (By similarity).  
 CC -!- CATALYTIC ACTIVITY: RH + reduced flavoprotein + O(2) = ROH +  
 CC oxidized flavoprotein + H(2)O.  
 CC -!- SUBCELLULAR LOCATION: Membrane-bound. Endoplasmic reticulum  
 CC (Potential).  
 CC -!- SIMILARITY: Belongs to the cytochrome P450 family.  
 CC  
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EMBL; AB003837; AAF59092.1; --  
 DR EMBL; AY061058; AAL28606.1; --  
 DR HSSP; P14779; 13PZ.  
 DR Flybase; FBgn0033292; Cyp4ad1.  
 DR InterPro; IPR001128; Cytochrome\_P450.  
 DR Pfam; PF00067; P450; 1.  
 DR PRINTS; PR00385; P450.  
 DR PROSITE; PS00086; CYTOCHROME\_P450; 1.  
 KW Oxidoreductase; Monooxygenase; Membrane; Heme; Microsome;  
 KW Endoplasmic reticulum; Hypothetical protein.  
 FT METAL 445 445 IRON (HEME AXIAL LIGAND) (BY SIMILARITY).  
 SQ SEQUENCE 516 AA; 58870 MW; 648EA22492AF58C7 CRC64;

Query Match 58.5%; Score 38; DB 1; Length 516;  
 Best Local Similarity 80.0%; Pred. No. 69;  
 Matches 8; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 LPVAVGLSPG 10  
 DB 469 LPVAVGLSPG 478  
 |||||

RESULT 14  
 EFG HAEIN STANDARD; PRT; 699 AA.  
 ID EFG HAEIN STANDARD; PRT; 699 AA.  
 AC P43925;  
 DT 01-NOV-1995 (Rel. 32, Created)  
 DT 01-NOV-1995 (Rel. 32, Last sequence update)  
 DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 DE Elongation factor G (EF-G).  
 GN FUSA OR FUS OR HI0579.

```

OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Haemophilus.
OX NCBI_TaxID=727;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=RG / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Utterback T.R., Hanna M.C., Spriggs T., Saudek D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghegan N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RT "Whole-genome random sequencing and assembly of Haemophilus influenzae
RT Rd.";
RL Science 269:496-512(1995).
CC -!- FUNCTION: This protein promotes the GTP-dependent translocation of
CC the nascent protein chain from the A-site to the P-site of the
CC ribosome.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
CC -!- SIMILARITY: Belongs to the GTP-binding elongation factor family.
CC EF-G/EF-2 subfamily.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC or send an email to license@isb-sib.ch).
CC
DR EMBL; U32739; AAC22237.1; -.
DR PIR; F64078; F64078.
DR HSSP; F13551; IELO.
DR TIGR; H10579; -.
DR HAVAP; MF_00054; -.
DR InterPro; IPR004540; EF-G.
DR InterPro; IPR000795; EF_GTPbind.
DR InterPro; IPR000640; EF_G.
DR InterPro; IPR009022; EFG_III_V.
DR InterPro; IPR005517; EFG_IV.
DR InterPro; IPR004161; EFTU_D2.
DR InterPro; IPR005225; Small_GTP.
DR InterPro; IPR009000; Translat_factor.
DR Pfam; PF00679; EF-G; 1.
DR Pfam; PF03764; EF-G; 1.
DR Pfam; PF00009; GTP_EFTU; 1.
DR Pfam; PF03144; GTP_EFTU_D2; 1.
DR PRINTS; PR00315; ELONGATNFCT.
DR TIGRFAMS; TIGR00484; EF-G; 1.
DR TIGRFAMS; TIGR00231; small_GTP; 1.
DR PROSITE; PS00301; EFATOR_GTP; 1.
KW Elongation factor; Protein biosynthesis; GTP-binding;
KW Complete proteome.
FT INIT MET 0 0 BY SIMILARITY.
FT NP_BIND 16 23 GTP (BY SIMILARITY).
FT NP_BIND 87 91 GTP (BY SIMILARITY).
FT NP_BIND 141 144 GTP (BY SIMILARITY).
SQ SEQUENCE 699 AA; 77132 MW; FBBAD39C0FG62801 CRC64;

Query Match 58.5%; Score 38; DB 1; Length 699;
Best Local Similarity 46.2%; Pred. No. 93;
Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPAVGLSPGEQE 13
Db 291 IPAIKGINPDETE 303

```

```

RESULT 15
EFG_PASMU STANDARD; PRT; 700 AA.
AC P57938;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Elongation factor G (EF-G).
GN FUSA OR PM1356.
OS Pasteurella multocida.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Pasteurella.
OX NCBI_TaxID=747;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Pm70;
RX MEDLINE=21145866; PubMed=11248100;
RA May B.J., Zhang Q., Li L.L., Paustian M.L., Whittam T.S., Kapur V.;
RA "Complete genomic sequence of Pasteurella multocida Pm70.";
RA Proc. Natl. Acad. Sci. U.S.A. 98:3460-3465(2001).
CC -!- FUNCTION: This protein promotes the GTP-dependent translocation of
CC the nascent protein chain from the A-site to the P-site of the
CC ribosome.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
CC -!- SIMILARITY: Belongs to the GTP-binding elongation factor family.
CC EF-G/EF-2 subfamily.
CC
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CC or send an email to license@isb-sib.ch).
CC
DR EMBL; AE006173; AAK03440.1; -.
DR HSSP; P13551; IELO.
DR HAVAP; MF_00054; -.
DR InterPro; IPR004540; EF-G.
DR InterPro; IPR000795; EF_GTPbind.
DR InterPro; IPR000640; EF_G.
DR InterPro; IPR009022; EFG_III_V.
DR InterPro; IPR005517; EFG_IV.
DR InterPro; IPR004161; EFTU_D2.
DR InterPro; IPR005225; Small_GTP.
DR InterPro; IPR009000; Translat_factor.
DR Pfam; PF00679; EF-G; 1.
DR Pfam; PF03764; EF-G; 1.
DR Pfam; PF00009; GTP_EFTU; 1.
DR Pfam; PF03144; GTP_EFTU_D2; 1.
DR PRINTS; PR00315; ELONGATNFCT.
DR TIGRFAMS; TIGR00484; EF-G; 1.
DR TIGRFAMS; TIGR00231; small_GTP; 1.
DR PROSITE; PS00301; EFATOR_GTP; 1.
KW Elongation factor; Protein biosynthesis; GTP-binding;
KW Complete proteome.
FT NP_BIND 16 23 GTP (BY SIMILARITY).
FT NP_BIND 87 91 GTP (BY SIMILARITY).
FT NP_BIND 141 144 GTP (BY SIMILARITY).
SQ SEQUENCE 700 AA; 77186 MW; 6CC161F7F9FA9C72 CRC64;

Query Match 58.5%; Score 38; DB 1; Length 700;
Best Local Similarity 46.2%; Pred. No. 93;
Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPAVGLSPGEQE 13
Db 292 IPAIKGINPDETE 304

```

Search completed: May 7, 2004, 12:34:33  
Job time: 4.68 secs

Result No.	Score	Query		Length	DB	ID	Description
		Match	%				
1	51	39.2	426	2	D72635	hypothetical prote	
2	50.5	38.8	615	2	AC3554	galactonate dehydr	
3	50	38.5	277	2	F82130	formyltetrahydrofo	
4	50	38.5	381	2	F75270	cytochrome P450 -	
5	49.5	38.1	603	2	A22952	dihydroxy-p acid de	
6	49.5	38.1	603	2	H98330	hypothetical prote	
7	49	37.7	118	2	S38491	Ig heavy chain - h	
8	49	37.7	278	2	E64131	formyltetrahydrofo	
9	48.5	37.3	559	2	S82503	inorganic phosphat	
10	48	36.9	522	2	T44369	pyruvate, water di	
11	48	36.9	567	2	AF0274	ribulokinase (EC 2	
12	48	36.9	780	2	B27561	phosphoenolpyruvat	
13	48	36.9	792	2	B82752	penicillin binding	
14	47	36.2	280	2	F90845	formyltetrahydrofo	
15	47	36.2	280	2	E85703	hypothetical prote	
16	47	36.2	280	2	C36871	formyltetrahydrofo	
17	47	36.2	280	2	AF0649	formyltetrahydrofo	
18	47	36.2	293	2	E71819	formyltetrahydrofo	
19	47	36.2	293	2	B64699	formyltetrahydrofo	
20	47	36.2	661	2	G84511	formyltetrahydrofo	
21	47	36.2	2109	1	I50421	hypothetical prote	
22	46.5	35.8	178	2	G82977	aggreacan precursor	
23	46.5	35.8	563	2	T36580	hypothetical prote	
24	46.5	35.8	656	2	B81692	hypothetical prote	
25	46	35.4	192	2	A82545	penicillin-binding	
26	46	35.4	271	2	C92841	hypothetical prote	
27	46	35.4	274	2	F81350	cysteine proteinas	
28	46	35.4	287	2	B70871	formyltetrahydrofo	
29	46	35.4	300	2	C69857	hypothetical prote	
			287	2	C69857	formyltetrahydrofo	

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Query Match      38.8%; Score 50.5; DB 2; Length 615;
Best Local Similarity 45.8%; Pred. No. 26;
Matches 11; Conservative 2; Mismatches 6; Indels 5; Gaps 1;

QY 4 LPAVVGSLP-----GQEQYHGGV 22
Db 324 IPLLNVNLPAGEYLGEDYHAGGV 347

RESULT 3
F82130
C:Species: Vibrio cholerae
C>Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
C:Accession: F82130
R:Heidelberger, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.J.;
I, R.K.; Mekalanos, J.J.; Venter, J.C.; Qin, H.; Dragoi, I.; Sellers, H.
Nature 406, 477-483, 2000
A:Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.
A:Reference number: AB2035; MUID:20406833; PMID:10952301
A:Accession: F82130
A>Status: Preliminary
A:Molecule type: DNA
A:Residues: 1-277 <HEI>
A:Cross-references: GB:AE003852; NID:9956533; PIDN:AAF95140.1; GSPDB:GN001
A:Experimental source: serogroup O1; strain N16961; biotype El Tor
C:Genetics:
A:Gene: VC1992
A:Map position: 1
C:Superfamily: phosphoribosylglycinamide formyltransferase; phosphoribosylglycinamide fo

Query Match      38.5%; Score 50; DB 2; Length 277;
Best Local Similarity 52.9%; Pred. No. 13;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPAVVGSLSPGQEQYHGG 20
Db 190 LPAFTGAKPYQAVERG 206

RESULT 4
F75270
Cytochrome P450 - Deinococcus radiodurans (strain R1)
C:Species: Deinococcus radiodurans
C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 17-Mar-2000
C:Accession: F75270
R:White, O.; Eisen, J.A.; Heidelberger, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.
Science 286, 1571-1577, 1999
A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.
A:Reference number: A75250; MUID:20036896; PMID:10567266
A:Accession: F75270
A>Status: Preliminary
A:Molecule type: DNA
A:Residues: 1-381 <WHI>
A:Cross-references: GB:AE002076; GB:AE000513; NID:96460285; PIDN:AAF12016.1; PID:9646029
A:Experimental source: strain R1
C:Genetics:
A:Gene: DR2473
A:Map position: 1

Query Match      38.5%; Score 50; DB 2; Length 381;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GLPAVVGSLP 12
Db 51 GLPAVVGSLP 60

```

```

RESULT 5
AB2952
dihydroxy-acid dehydratase [imported] - Agrobacterium tumefaciens (strain C58, Dupont)
C:Species: Agrobacterium tumefaciens
C>Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-Nov-2002
C:Accession: AB2952
R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo,
erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutyavin, T.; Levy, R.; Li, M.; McClell
; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,
ster, E.W.
A:Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
A:Reference number: AB2577; MUID:21608550; PMID:11743193
A:Accession: AB2952
A>Status: Preliminary
A:Molecule type: DNA
A:Residues: 1-603 <KUR>
A:Cross-references: GB:AE008689; PIDN:AAU44035.1; PID:GL7741597; GSPDB:GN00187
A:Experimental source: strain C58 (Dupont)
C:Genetics:
A:Gene: ilvD
A:Map position: linear chromosome
C:Superfamily: dihydroxy-acid dehydratase

Query Match      38.1%; Score 49.5; DB 2; Length 603;
Best Local Similarity 45.8%; Pred. No. 36;
Matches 11; Conservative 2; Mismatches 6; Indels 5; Gaps 1;

QY 4 LPAVVGSLP-----GQEQYHGGV 22
Db 312 VPLLNVNLPAGEYLGEDYHAGGV 335

RESULT 6
H98330
hypothetical protein AGR_L_3190 [imported] - Agrobacterium tumefaciens (strain C58, Cer
C:Species: Agrobacterium tumefaciens
C>Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 18-Nov-2002
C:Accession: H98330
R:Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Ouello, B.; Goldman
A.; Liu, F.; Wollam, C.; Allinger, M.; Doughty, D.; Scott, C.; Lappas, C.; Markelz, B.
Science 294, 2323-2328, 2001
A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tu
A:Reference number: A97359; MUID:21608551; PMID:11743194
A:Accession: H98330
A>Status: Preliminary
A:Molecule type: DNA
A:Residues: 1-603 <KUR>
A:Cross-references: GB:AE007870; PIDN:AAK90170.1; PID:GL5160173; GSPDB:GN00170
C:Genetics:
A:Gene: AGR_L_3190
A:Map position: linear chromosome
C:Superfamily: dihydroxy-acid dehydratase

Query Match      38.1%; Score 49.5; DB 2; Length 603;
Best Local Similarity 45.8%; Pred. No. 36;
Matches 11; Conservative 2; Mismatches 6; Indels 5; Gaps 1;

QY 4 LPAVVGSLP-----GQEQYHGGV 22
Db 312 VPLLNVNLPAGEYLGEDYHAGGV 335

RESULT 7
S38491
IG heavy chain - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 23-Jul-1999
C:Accession: S38491
R:Markes, J.D.; Ouweland, W.H.; Bye, J.M.; Finnern, R.; Gorick, B.D.; Voak, D.; Thorpe,
submitted to the EMBL Data Library, June 1993
A:Description: Human antibody fragments specific for human blood group antigens from a

```

A;Reference number: S38488

A;Accession: S38491

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-118 <MAR>

A;Cross-references: EMBL:Z23032; NID:G414029; PIDN:CAA80567.1; PID:G414030

A;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotrimer; immunoglobulin

F;15-97/Domain: immunoglobulin homology <IMV>

Query Match 37.7%; Score 49; DB 2; Length 118;  
Best Local Similarity 50.0%; Pred. No. 7.3;  
Matches 10; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

QY 3 GLPAVVGSLSPGQYHRGV 22

Db 44 GLEWVGIGPGGDTYFGSV 53

RESULT 8

E64131

formyltetrahydrofolate deformylase (EC 3.5.1.10) - Haemophilus influenzae (strain Rd KW2)

C;Species: Haemophilus influenzae

C;Date: 18-Aug-1995 #sequence\_revision 18-Aug-1995 #text\_change 17-Mar-2000

C;Accession: E64131; FN0606

R;Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A.

; Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.W.; Weidman, J.

; D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghagen, N.S.M.

Science 269, 496-512, 1995

A;Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,

A;Title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.

A;Reference number: A64000; MUID:95350630; PMID:7542800

A;Accession: E64131

A;Status: nucleic acid sequence not shown; translation not shown

A;Molecule type: DNA

A;Residues: 1-278 <TYGR>

A;Cross-references: GB:U32833; GB:I42023; NID:G1574432; PIDN:AAC23236.1; PID:G1574433; T

R;Maskell, D.

Gene 129, 155-156, 1993

A;Title: Cloning and sequencing of the Haemophilus influenzae aroA gene.

A;Reference number: JN0758; MUID:93328119; PMID:8335255

A;Accession: FN0606

A;Molecule type: DNA

A;Residues: 64-114, 'FNR', 118-137, 'PK', 141-204, 'E', 206-278 <MAS>

A;Cross-references: GB:L04686; NID:G148863; PIDN:AAA24942.1; PID:G148864

C;Genetics:

A;Gene: purN

C;Function:

A;Description: catalyzes hydrolysis of 10-formyltetrahydrofolate to tetrahydrofolate and

A;Pathway: one-carbon metabolism

A;Note: activated by methionine and inhibited by glycine

C;Superfamily: phosphoribosylglycinamide formyltransferase; phosphoribosylglycinamide fo

C;Keywords: hydrolase

F;85-277/Domain: phosphoribosylglycinamide formyltransferase homology <PRGF>

F;223/Active site: Asp #status Predicted

Query Match 37.7%; Score 49; DB 2; Length 278;

Best Local Similarity 52.9%; Pred. No. 18;

Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPAVGLSPGQYHRG 20

Db 191 LPAFIGAKPYQAYKRG 207

RESULT 9

S62503

inorganic phosphate transporter - fission yeast (Schizosaccharomyces pombe)

C;Species: Schizosaccharomyces pombe

C;Date: 16-May-1996 #sequence\_revision 13-Mar-1997 #text\_change 10-Dec-1999

C;Accession: T38287; S62503

R;Niblett, D.; Harris, D.; Barrell, B.G.; Rajandream, M.A.; Walsh, S.V.

submitted to the EMBL Data Library, October 1995

A;Reference number: Z21783

A;Accession: T38287

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-559 <NI2>

A;Cross-references: EMBL:Z64354; NID:G1039338; PIDN:CAA91247.1; PID:G1039350; GSPDB:GNO

A;Experimental source: strain 972h-; cosmid C23D3

C;Genetics:

A;Gene: SPDB:SPAC23D3.12

A;Map position: 1R

C;Superfamily: probable inorganic phosphate transport protein PHO84

Query Match 37.3%; Score 48.5; DB 2; Length 559;

Best Local Similarity 47.6%; Pred. No. 46;

Matches 10; Conservative 3; Mismatches 5; Indels 3; Gaps 1;

QY 8 VQLSPGQYH---RGVGV 25

Db 365 IGFSGKNEYHTLRGAIGNL 385

RESULT 10

T44369

Pyruvate, water dikinase (EC 2.7.9.2) [imported] - Deinococcus radiodurans (fragment)

C;Species: Deinococcus radiodurans

C;Date: 21-Jan-2000 #sequence\_revision 21-Jan-2000 #text\_change 03-Jun-2002

C;Accession: T44369

R;Narumi, I.; Islam, S.; Cherdchu, K.; Kikuchi, M.; Watanabe, H.; Kitayama, S.; Yamamoto

submitted to the EMBL Data Library, August 1998

A;Description: I88301: the second insertion sequence element from Deinococcus radiodura

A;Reference number: Z22755

A;Accession: T44369

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-522 <NAR>

A;Cross-references: EMBL:AB016803; PIDN:BA432387.1

A;Experimental source: strain KD8301

C;Genetics:

A;Note: ppsA

C;Superfamily: Escherichia coli pyruvate, water dikinase; phosphotransferase system enzy

C;Keywords: transferase

Query Match 36.9%; Score 48; DB 2; Length 522;

Best Local Similarity 50.0%; Pred. No. 50;

Matches 9; Conservative 2; Mismatches 7; Indels 0; Gaps 0;

QY 3 GLPAVVGSLSPGQYHRG 20

Db 160 GIPAVVGTCNATRELNG 177

RESULT 11

AF0274

ribulokinase (EC 2.7.1.16) [imported] - Yersinia pestis (strain CO92)

C;Species: Yersinia pestis

C;Date: 02-Nov-2001 #sequence\_revision 02-Nov-2001 #text\_change 27-Nov-2001

C;Accession: AF0274

R;Parkhill, J.; Wren, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.G.; Prentice, M.B

deno-Tarraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.;

il, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; Whitehead, S.; Barrell,

Nature 413, 523-527, 2001

A;Title: Genome sequence of Yersinia pestis, the causative agent of plague.

A;Reference number: AB0001; MUID:21470413; PMID:11586360

A;Accession: AF0274

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-567 <KUR>

A;Cross-references: GB:AL590842; PIDN:CAC91058.1; PID:G15980250; GSPDB:GN00175

C;Genetics:

A;Gene: araB

C;Superfamily: ribulokinase

C;Keywords: phosphotransferase





GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 9 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-5  
Perfect score: 130  
Sequence: 1 MAGLPAVGLSPGEQYHGGVGL 25

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	49	37.7	278	1	PURU_HAEIN
2	48.5	37.3	559	1	YABC_SCHPO
3	48	36.9	563	1	ARAB_YERPE
4	48	36.9	780	1	PEPSA_DEIRA
5	47	36.2	280	1	PURU_ECOL6
6	47	36.2	280	1	PURU_ECOLI
7	47	36.2	567	1	ARAB_VIBPA
8	47	36.2	813	1	CADM_MOUSE
9	47	36.2	2109	1	PGCA_CHICK
10	46	35.4	508	1	COBI_MYCTU
11	46	35.4	1172	1	CNA2_MOUSE
12	45	34.6	479	1	PTSB_VIBAL
13	45	34.6	961	1	ACON_MYCAV
14	44.5	34.2	618	1	ILVD_CHRVO
15	44.5	34.2	2054	1	M18A_HUMAN
16	44	33.8	151	1	SODC_XIPGL
17	44	33.8	433	1	ENOB_RABIT
18	44	33.8	565	1	ARAB_ECOL57
19	44	33.8	565	1	ARAB_ECOL6
20	44	33.8	565	1	ARAB_ECOLI
21	44	33.8	614	1	ARAB_SALTI
22	44	33.8	568	1	CPRI_DROME
23	44	33.8	617	1	ILVD_STRFO
24	44	33.8	658	1	VG18_BPT4
25	44	33.8	677	1	SG1_HUMAN
26	44	33.8	777	1	Li10_ADE41
27	44	33.8	813	1	CADM_RAT
28	44	33.8	1065	1	RPOB_MARPO
29	44	33.8	3664	1	MINT_HUMAN
30	44	33.8	4351	1	FAT2_RAT
31	43.5	33.5	173	1	RUVG_XYLFA
32	43.5	33.5	173	1	RUVG_XYLFT
33	43.5	33.5	968	1	CTDI_HUMAN

34	43	33.1	211	1	COBL_METJA
35	43	33.1	325	1	CA19_RAT
36	43	33.1	326	1	MER_MERTI
37	43	33.1	364	1	COAL_SV40
38	43	33.1	382	1	FETE_HUMAN
39	43	33.1	390	1	COBL_MYCTU
40	43	33.1	481	1	GLGA_RHILO
41	43	33.1	505	1	Y76J_CAEEL
42	43	33.1	516	1	C4AD_DROME
43	43	33.1	544	1	CH60_PROAC
44	43	33.1	561	1	LCFA_SALTY
45	43	33.1	628	1	HNFA_MOUSE

O58917	methanococc
P20850	rattus norv
Q9uxp0	methanolobu
P03087	simian viru
Q9ugm5	homo sapien
Q10671	mycobacteri
Q985p2	rhizobium l
P90938	caenorhabdi
Q9v4t3	drosophila
Q9k2u4	propionibac
Q8xg98	salmonella
P22361	mus musculu

#### ALIGNMENTS

RESULT 1  
PURU\_HAEIN STANDARD; PRT; 278 AA.  
AC Q03432;  
DT 01-OCT-1994 (Rel. 30, Created)  
DT 01-NOV-1995 (Rel. 32, Last sequence update)  
DT 28-FEB-2003 (Rel. 41, Last annotation update)  
DE Formyltetrahydrofolate deformylase (RC 3.5.1.10) (Formyl-FH(4) hydrolase).  
DE hydrolase).  
GN PURU OR H11588.  
OS Haemophilus influenzae.  
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;  
OC Pasteurellaceae; Haemophilus.  
OX NCBI\_TaxID=727;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=Rd / KW20 / ATCC 51907;  
RX MEDLINE=95350630; PubMed=7542800;  
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F., Kierlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M., McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D., Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M., Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D., Uterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C., Fine L.D., Fritchman J.L., Fuhmann J.L., Geoghagen N.S.M., Ghem C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O., Venter J.C.;  
RL "Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.";  
RL Science 269:496-512(1995).  
RN [2]  
RP SEQUENCE OF 64-278 FROM N.A.  
RC STRAIN=RM 7004 / Serotype B;  
RX MEDLINE=93328119; PubMed=8335255;  
RA Maskell D.J.;  
RL "Cloning and sequencing of the Haemophilus influenzae aroA gene.";  
RL Gene 129:153-156(1993).

CC -!- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF 5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL (BY SIMILARITY).  
CC -!- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate + tetrahydrofolate.  
CC -!- ENZYME REGULATION: Activated by methionine, inhibited by glycine (By similarity).  
CC -!- PATHWAY: De novo purine biosynthesis.  
CC -!- SUBUNIT: Homohexamer (By similarity).  
CC -!- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).  
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CC -----
DR EMBL; U22833; AAC23236.1; -.
DR EMBL; L04686; AAA24942.1; -.
DR PIR; E64131; E64131.
DR HSP; P08179; IGRC.
DR TIGR; H11588; -.
DR InterPro; IPR002912; ACT.
DR InterPro; IPR002376; formyl_transf.
DR InterPro; IPR004810; Pur.
DR Pfam; PF01842; ACT; 1.
DR Pfam; PF00551; formyl_transf; 1.
DR PRINTS; PR01575; FFH4HYDLASE.
DR TIGRfams; TIGR00655; Pur; 1.
KW Purine biosynthesis; Hydrolase; One-carbon metabolism;
KW Complete proteome.
FT ACT_SITE 223 BY SIMILARITY.
FT CONFLICT 115 117 VIG -> RNR (IN REF. 2).
FT CONFLICT 138 140 HEN -> PK (IN REF. 2).
FT CONFLICT 205 205 K -> E (IN REF. 2).
SQ SEQUENCE 278 AA; 32173 MW; 7F375AB3C422EC4B CRC64;

Query Match 37.7%; Score 49; DB 1; Length 278;
Best Local Similarity 52.9%; Pred. No. 8.6;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPVVGLSPGGEQYHRG 20
DB 191 LPAFIGAKPYQAYKRG 207

RESULT 2
YAEC SCHPO STANDARD; PRT; 559 AA.
AC Q09852;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DE Putative inorganic phosphate transporter C3D3.12.
GN SPAC2D3.12.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OX NCBI_TaxID=4896;
RN [1]
RC SEQUENCE FROM N.A.
RE STRAIN=972;
RX MEDLINE=21848401; PubMed=11859360;
RA Wood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgouros J., Peat N., Hayes J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Voicakeert G., Aert R., Robben J., Grymonprez B.,
RA Welljens I., Vansteels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Moestl D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Carrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;

```

The genome sequence of *Schizosaccharomyces pombe*.  
 Nature 415:871-880(2002).  
 -!- FUNCTION: High-affinity transporter for external inorganic phosphate (By similarity).  
 -!- SUBCELLULAR LOCATION: Integral membrane protein (Probable).  
 -!- SIMILARITY: BELONGS TO THE SUGAR TRANSPORTER FAMILY. STRONG, TO YEAST PHO84.

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EMBL; Z64354; CAA91247.1; -.  
 PIR; T38287; S62503.  
 GenBank; SPombe; SPAC2D3.12; -.  
 InterPro; IPR007114; MFS.  
 InterPro; IPR004738; Phos\_permease.  
 InterPro; IPR005828; Sub\_transporter.  
 InterPro; IPR005829; Sug\_transporter.  
 Pfam; PF00083; sugar\_tr; 1.  
 TIGRfams; TIGR00887; 2A0109; 1.  
 PROSITE; PS00850; MFS; 1.  
 PROSITE; PS00216; SUGAR\_TRANSPORT\_1; 1.  
 PROSITE; PS00217; SUGAR\_TRANSPORT\_2; 1.  
 KW Hypothetical protein; Phosphate transport; Transport; Transmembrane.  
 FT DOMAIN 1 45 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 46 66 1 (POTENTIAL).  
 FT DOMAIN 67 94 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 95 115 2 (POTENTIAL).  
 FT DOMAIN 116 118 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 119 139 3 (POTENTIAL).  
 FT DOMAIN 140 144 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 145 165 4 (POTENTIAL).  
 FT DOMAIN 166 188 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 189 209 5 (POTENTIAL).  
 FT DOMAIN 210 224 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 225 245 6 (POTENTIAL).  
 FT DOMAIN 246 335 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 336 356 7 (POTENTIAL).  
 FT DOMAIN 357 381 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 382 402 8 (POTENTIAL).  
 FT DOMAIN 403 409 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 410 430 9 (POTENTIAL).  
 FT DOMAIN 431 438 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 439 459 10 (POTENTIAL).  
 FT DOMAIN 460 474 CYTOPLASMIC (POTENTIAL).  
 FT TRANSMEM 475 495 11 (POTENTIAL).  
 FT DOMAIN 496 497 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 498 518 12 (POTENTIAL).  
 FT DOMAIN 519 559 CYTOPLASMIC (POTENTIAL).  
 SQ SEQUENCE 559 AA; 62316 MW; 2E8958F86C2092E2 CRC64;

Query Match 37.3%; Score 48.5; DB 1; Length 559;  
 Best Local Similarity 47.6%; Pred. No. 21;  
 Matches 10; Conservative 3; Mismatches 5; Indels 3; Gaps 1;

QY 8 VGLSPGEQYH---RGVGVL 25  
 DB 365 IGFSGKNEYHTLMRGAINL 385

RESULT 3  
 ARAB\_YERPE STANDARD; PRT; 563 AA.  
 ID ARAB\_YERPE  
 AC P58543;  
 DT 28-FEB-2003 (Rel. 41, Created)  
 DT 28-FEB-2003 (Rel. 41, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE L-ribulokinase (EC 2.7.1.16).

```

Query Match          36.9%;   Score 48;   DB 1;   Length 780;
Best Local Similarity 50.0%;   Pred. No. 35;
Matches 9; Conservative 2; Mismatches 7; Indels 0; Gaps 0;

QY      3  GLPAVVGLSGGEQYHRG  20
      ||:|||||
      :|||
Db      418  GIPAVVGIGNATRELHG  435

```



RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,  
 RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,  
 RA Mau B., Shao Y.;  
 RT "The complete genome sequence of *Escherichia coli* K-12.";  
 RL Science 277:1453-1474 (1997).  
 RN [4]  
 RN SEQUENCE FROM N.A.  
 RC STRAIN=K12;  
 RX MEDLINE=97061202; PubMed=8905232;  
 RA Oshima T., Aiba H., Baba T., Fujita K., Hayashi K., Honjo A.,  
 RA Ikemoto K., Inada T., Itoh T., Kajihara M., Kanai K., Kashimoto K.,  
 RA Kimura S., Kitagawa M., Makino K., Masuda S., Miki T., Mizobuchi K.,  
 RA Mori H., Motomura K., Nakamura Y., Nishimoto H., Nishio Y., Saito N.,  
 RA Sampei G., Seki Y., Tagami H., Takemoto K., Wada C., Yamamoto Y.,  
 RA Yano M., Horiiuchi T.;  
 RT "A 718-Kb DNA sequence of the *Escherichia coli* K-12 genome  
 corresponding to the 12.7-28.0 min region on the linkage map.";  
 RL DNA Res. 3:137-155 (1996).  
 RN [5]  
 RN CHARACTERIZATION.  
 RP MEDLINE=95173107; PubMed=7868604;  
 RA Nagy P.J., Marolewski A., Benkovic S.J., Zalkin H.;  
 RT "Formyltetrahydrofolate hydrolase, a regulatory enzyme that functions  
 to balance pools of tetrahydrofolate and one-carbon tetrahydrofolate  
 adducts in *Escherichia coli*.";  
 RL J. Bacteriol. 177:1292-1298 (1995).  
 CC -!- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES  
 CC THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF  
 CC 5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC  
 CC GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL.  
 CC -!- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate +  
 CC tetrahydrofolate.  
 CC -!- ENZYME REGULATION: Activated by methionine, inhibited by glycine.  
 CC -!- PATHWAY: De novo purine biosynthesis.  
 CC -!- SUBUNIT: Homohexamer.  
 CC -!- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).  
 CC -----  
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 CC -----  
 DR EMBL; L20251; AAC36846.1; -;  
 DR EMBL; M64675; AAA16860.1; ALT INIT.  
 DR EMBL; AE000221; AAC74314.1; -;  
 DR EMBL; D90758; BAA36100.1; -;  
 DR EMBL; D90759; BAA36112.1; -;  
 DR EMBL; D90852; BAA16026.1; -;  
 DR PIR; C36871; C36871.  
 DR EcGene; EG11819; purU.  
 DR InterPro; IPR002912; ACT.  
 DR InterPro; IPR002376; formyl\_transf.  
 DR InterPro; IPR004810; PurU.  
 DR Pfam; PF01842; ACT; 1.  
 DR Pfam; PF00551; formyl\_transf; 1.  
 DR PRINTS; PR01575; FFH4HYDLRLASE.  
 DR TIGRFAMS; TIGR00655; PurU; 1.  
 KW Purine biosynthesis; Hydrolase; One-carbon metabolism;  
 KW Complete proteome.  
 FT ACT SITE 225 225 BY SIMILARITY.  
 SQ SEQUENCE 280 AA; 31934 MW; 5667406D2727A2C2 CRC64;

Query Match 36.2%; Score 47; DB 1; Length 280;  
 Best Local Similarity 52.9%; Pred. No. 17;  
 Matches 9; Conservative 1; Mismatches 7; Indels 0; Gaps 0;

QY 4 LPVAVGLSPGQEQYHRG 20  
 DB 193 LPAFIGARPHQAYERG 209

RESULT 7  
 ARAB\_VIBPA STANDARD; PRT; 567 AA.  
 ID ARAB\_VIBPA  
 AC Q87FK5;  
 DT 10-OCT-2003 (Rel. 42, Created)  
 DT 10-OCT-2003 (Rel. 42, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE L-ribulokinase (EC 2.7.1.16).  
 GN ARAB OR VPA1674.  
 OS Vibrio parahaemolyticus.  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Vibrionales;  
 OC Vibrionaceae; Vibrio.  
 OX NCBI\_TaxID=670;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RP STRAIN=RMD 2210633 / Serotype O3:K6;  
 RX MEDLINE=22508454; PubMed=12620739;  
 RA Makino K., Oshima K., Kurokawa K., Yokoyama K., Uda T., Tagomori K.,  
 RA Iijima Y., Najima M., Nakano M., Yamashita A., Kubota Y., Kimura S.,  
 RA Yasunaga T., Honda T., Shinagawa H., Hattori M., Iida T.;  
 RT "Genome sequence of *Vibrio parahaemolyticus*: a pathogenic mechanism  
 distinct from that of *V. cholerae*.";  
 RL Lancet 361:743-749 (2003).  
 CC -!- CATALYTIC ACTIVITY: ATP + L-ribulose = ADP + L-ribulose 5-  
 CC phosphate.  
 CC -!- PATHWAY: L-arabinose catabolism; second step.  
 CC -!- SIMILARITY: Belongs to the ribulokinase family.  
 CC -----  
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 CC -----  
 DR EMBL; AP005089; BAC63017.1; -;  
 DR HAMAP; MF\_00520; -; 1.  
 DR InterPro; IPR000577; FGGY\_kin.  
 DR Pfam; PF00370; FGGY; 1.  
 DR Pfam; PF02782; FGGY\_C; 1.  
 DR Pfam; PF02782; Kinase; Arabinose catabolism: Complete proteome.  
 KW Transferase; Kinase; Arabinose catabolism: Complete proteome.  
 SQ SEQUENCE 567 AA; 61788 MW; B303B547EBEA9B8B CRC64;  
 -----  
 Query Match 36.2%; Score 47; DB 1; Length 567;  
 Best Local Similarity 42.9%; Pred. No. 35;  
 Matches 9; Conservative 4; Mismatches 8; Indels 0; Gaps 0;  
 -----  
 QY 3 GLPAAVGLSPGQEQYHRG 23  
 DB 267 GLPEGTAIGEFDCMGAVG 287  
 -----  
 RESULT 8  
 CADM\_MOUSE STANDARD; PRT; 813 AA.  
 ID CADM\_MOUSE  
 AC Q8WT5;  
 DT 28-FEB-2003 (Rel. 41, Created)  
 DT 28-FEB-2003 (Rel. 41, Last sequence update)  
 DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 DE Cadherin-22 precursor (PB-cadherin).  
 GN CDH22.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_TaxID=10090;  
 RN [1]  
 RN SEQUENCE FROM N.A., AND DEVELOPMENTAL STAGE.  
 RC STRAIN=ICR; TISSUE=Brain;  
 RX MEDLINE=99326347; PubMed=10398531;  
 RA Kitajima K., Koshimizu U., Nakamura T.;  
 RT "Expression of a novel type of classic cadherin, PB-cadherin in

```

RT developing brain and limb buds. ";
RL Dev. Dyn. 215:206-214(1999).
CC
CC -!- FUNCTION: Cadherins are calcium dependent cell adhesion proteins.
CC They preferentially interact with themselves in a homophilic
CC manner in connecting cells; cadherins may thus contribute to the
CC sorting of heterogeneous cell types. P3-cadherins may have a role
CC in the morphological organization of pituitary gland and brain
CC tissues.
CC
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC
CC -!- TISSUE SPECIFICITY: Predominantly expressed in brain. Abundant in
CC olfactory bulb, cerebellum, and cerebellum, less in pons, medulla,
CC and spinal cord. Low expression in heart. No expression in lung,
CC liver, spleen, kidney, testis, stomach, intestine, colon, and
CC placenta.
CC
CC -!- DEVELOPMENTAL STAGE: Expressed at 9.5 dpc onwards. At 10.5 dpc, in
CC brain (telencephalic vesicles and isthmus), spinal cord and limb
CC buds (in the zone of polarizing activity). At 14.5 dpc, in
CC olfactory bulb and cerebellum.
CC
CC -!- INDUCTION: Down-regulated by thyroid hormone.
CC
CC -!- SIMILARITY: Contains 5 cadherin domains.
CC
CC -----
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CC
CC -----
DR EMBL; AB019618; BAA34426.1; -.
DR MGD; MGI:1341843; Cdh22.
DR HSP; P15116; INCU.
DR InterPro: IPR002126; Cadherin.
DR InterPro: IPR000233; Cadherin_C term.
DR Pfam; PF00028; cadherin; 5.
DR Pfam; PF01049; Cadherin_C term; 1.
DR PRINTS; PR00205; CADHERIN.
DR SMART; SM00112; CA; 5.
DR PROSITE; PS00232; CADHERIN_1; 2.
DR PROSITE; PS00268; CADHERIN_2; 5.
DR Cell adhesion; Glycoprotein; Transmembrane; Calcium-binding; Repeat;
KW Signal.
FT SIGNAL. 1 33
FT CHAIN 34 813
FT DOMAIN 33 621
FT TRANSMEM 622 642
FT DOMAIN 643 813
FT DOMAIN 61 165
FT DOMAIN 166 274
FT DOMAIN 275 391
FT DOMAIN 392 495
FT DOMAIN 496 613
FT CARBOHYD 159 159
FT CARBOHYD 463 463
FT CARBOHYD 609 609
SQ SEQUENCE 813 AA; 88021 MW; 5510F9848D976567 CRC64;

Query Match
Best Local Similarity 36.2%; Score 47; DB 1; Length 813;
Matches 11; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 2 AGLPAVGLSPGEQYHRRGVG 23
Db 34 ASTPAPSSLSPGAQDNQLGAG 55

RESULT 9
PGCA CHICK
ID PGCA CHICK STANDARD; PRT; 2109 AA.
AC P07898; Q90820; Q90991; Q91047;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)

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DE DE Aggrecan core protein precursor (Cartilage-specific proteoglycan core
GN GN protein) (CSPCP).
OS AGC1.
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=White leghorn; TISSUE=Embryo;
RX MEDLINE=94043149; PubMed=8226878;
RA Li H., Schwartz N.B., Vertel B.M.;
RT "cDNA cloning of chick cartilage chondroitin sulfate (aggrecan) core
RT protein and identification of a stop codon in the aggrecan gene
RT associated with the chondrodystrophy, nanomelia."
RL J. Biol. Chem. 268:23504-23511(1993).
RN [2]
RP SEQUENCE OF 1042-1559 FROM N.A.
RC TISSUE=Embryo;
RX MEDLINE=90307744; PubMed=1694853;
RA Krueger R.C. Jr., Fields T.A., Mensch J.R. Jr., Schwartz N.B.;
RT "Chick cartilage chondroitin sulfate proteoglycan core protein. II.
RT Nucleotide sequence of cDNA clone and localization of the S103L
RT epitope."
RL J. Biol. Chem. 265:12088-12097(1990).
RN [3]
RP SEQUENCE OF 1-1855 AND 1893-2109 FROM N.A.
RC TISSUE=Cartilage;
RX MEDLINE=93111968; PubMed=1339285;
RA Chandrasekaran L., Tanzer M.L.;
RT "Molecular cloning of chicken aggrecan. Structural analyses."
RL Biochem. J. 288:903-910(1992).
RN [4]
RP ERRATUM.
RX MEDLINE=94107258; PubMed=8280087;
RA Chandrasekaran L., Tanzer M.L.;
RL Biochem. J. 296:885-887(1993).
RN [5]
RP SEQUENCE OF 1492-1610 FROM N.A.
RC STRAIN=White leghorn; TISSUE=Chondrocytes;
RX MEDLINE=95129519; PubMed=7827752;
RA Primorac D., Stover M.L., Clark S.H., Rowe D.W.;
RT "Molecular basis of nanomelia, a heritable chondrodystrophy of
RT chicken."
RL Matrix Biol. 14:297-305(1994).
RN [6]
RP SEQUENCE OF 1894-2109 FROM N.A.
RX MEDLINE=89008500; PubMed=3170613;
RA Tanaka T., Har-El R., Tanzer M.L.;
RT "Partial structure of the gene for chicken cartilage proteoglycan
RT core protein."
RL J. Biol. Chem. 263:15831-15835(1988).
RN [7]
RP SEQUENCE OF 1693-1855 AND 1893-2109 FROM N.A.
RX MEDLINE=86259736; PubMed=3460082;
RA Sai S., Tanaka T., Koshier R.A., Tanzer M.L.;
RT "Cloning and sequence analysis of a partial cDNA for chicken
RT cartilage proteoglycan core protein."
RL Proc. Natl. Acad. Sci. U.S.A. 83:5081-5085(1986).
CC -!- FUNCTION: This proteoglycan is a major component of extracellular
CC matrix of cartilaginous tissues. A major function of this protein
CC is to resist compression in cartilage. It binds avidly to
CC hyaluronic acid via an amino-terminal globular region. May play a
CC regulatory role in the matrix assembly of the cartilage.
CC -!- SUBCELLULAR LOCATION: Secreted; extracellular matrix (By
CC similarity).
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1;
CC IsoId=P07898-1; Sequence=Displayed;
CC Name=2;
CC IsoId=P07898-2; Sequence=VSP_003073;

```

CC CC -!- DOMAIN: Two globular domains, G1 and G2, comprise the amino terminus of the proteoglycan, while another globular region, G3, makes up the COOH terminus. G1 contains link domains and thus consists of three disulfide-bonded loop structures designated as the A, B, B' motifs. G2 is similar to G1. The keratan sulfate (KS) and the chondroitin sulfate (CS) attachment domains lie between G2 and G3.

CC CC -!- PTM: Contains mostly chondroitin sulfate, but also keratan sulfate chains, N-linked and O-linked oligosaccharides.

CC CC -!- DISEASE: DEFECTS IN AGC1 ARE THE CAUSE OF NANOMELIA, A LETHAL CONNECTIVE TISSUE DISORDER AFFECTING CARTILAGE DEVELOPMENT (CHONDRODYSPLASIA) CHARACTERIZED BY SHORTENED AND MALFORMED LIMBS. AGGREGAN IS TRUNCATED AT ITS C-TERMINAL IN THE CS-2 BINDING DOMAIN AND IS NOT ANYMORE SECRETED FROM THE CHONDROCYTES.

CC CC -!- SIMILARITY: Belongs to the aggrecan/versican proteoglycan family.

CC CC -!- SIMILARITY: Contains 1 immunoglobulin-like V-type domain.

CC CC -!- SIMILARITY: Contains 4 link domains.

CC CC -!- SIMILARITY: Contains 1 EGF-like domain.

CC CC -!- SIMILARITY: Contains 1 C-type lectin family domain.

CC CC -!- SIMILARITY: Contains 1 Sushi (SCR) domain.

CC CC -----

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CC CC -----

DR EMBL; L21913; AAB19128.1; -;  
DR EMBL; M38187; AAA48731.1; -;  
DR EMBL; M88101; -; NOT ANNOTATED\_CDS.  
DR EMBL; S74657; AAC60751.1; -;  
DR EMBL; S74656; AAC60751.1; JOINED.  
DR EMBL; J04028; AAA48719.1; -;  
DR EMBL; M13993; AAA48720.1; -;  
DR PIR; I50421; I50421.  
DR HSP; P08709; IBF9.  
DR InterPro; IPR002353; AntifreezeII.  
DR InterPro; IPR000152; Asx hydroxyl\_s.  
DR InterPro; IPR000742; EGF\_2.  
DR InterPro; IPR001861; EGF\_Ca.  
DR InterPro; IPR006209; EGF\_Like.  
DR InterPro; IPR007110; Ig-like.  
DR InterPro; IPR003599; Ig.  
DR InterPro; IPR001304; Linkin\_C.  
DR InterPro; IPR000538; Link.  
DR InterPro; IPR003324; SGXSG.  
DR InterPro; IPR000436; Sushi\_SCR\_CCP.  
DR Pfam; PF00008; EGF; 1.  
DR Pfam; PF00047; Ig; 1.  
DR Pfam; PF00059; lectin\_c; 1.  
DR Pfam; PF02339; SGXSG; 56.  
DR Pfam; PF00084; sushi; 1.  
DR Pfam; PF00193; Xlink; 4.  
DR PRINTS; PR00356; ANTIFREEZEII.  
DR PRINTS; PR01265; LINKMODULE.  
DR ProDom; PD000918; Link; 4.  
DR SMART; SM00032; CCP; 1.  
DR SMART; SM00034; CLECT; 1.  
DR SMART; SM00179; EGF\_CA; 1.  
DR SMART; SM00409; Ig; 1.  
DR SMART; SM00445; LINK; 4.  
DR PROSITE; PS00010; ASX HYDROXYL; 1.  
DR PROSITE; PS00615; C-TYPE LECTIN\_1; 1.  
DR PROSITE; PS00441; C-TYPE LECTIN\_2; 1.  
DR PROSITE; PS00022; EGF\_1; 1.  
DR PROSITE; PS00026; EGF\_3; 1.  
DR PROSITE; PS01187; EGF\_CA; 1.  
DR PROSITE; PS00835; IG\_LIKE; 1.  
DR PROSITE; PS01241; LINK; 4.  
KW Glycoprotein; Proteoglycan; Lectin; Signal; Sushi; EGF-like domain;  
KW Alternative splicing; Repeat; Immunoglobulin domain.

FT	SIGNAL	1	16	POTENTIAL.
FT	CHAIN	17	2109	AGGREGAN CORE PROTEIN.
FT	DOMAIN	34	143	IG-LIKE V-TYPE.
FT	DOMAIN	166	243	LINK 1.
FT	DOMAIN	264	346	LINK 2.
FT	DOMAIN	537	614	LINK 3.
FT	DOMAIN	635	716	LINK 4.
FT	DOMAIN	1363	1742	19 X 20 AA TANDEM-REPEAT.
FT	DOMAIN	1855	1892	EGF-LIKE.
FT	DOMAIN	1901	2019	C-TYPE LECTIN.
FT	DOMAIN	2023	2081	SUSHI.
FT	DOMAIN	48	137	G1-A.
FT	DOMAIN	148	243	G1-B.
FT	DOMAIN	249	346	G1-B'.
FT	DOMAIN	519	613	G2-B.
FT	DOMAIN	620	715	G2-B'.
FT	DOMAIN	718	803	KS.
FT	DOMAIN	805	1264	CS-1.
FT	DOMAIN	1265	1742	CS-2.
FT	DOMAIN	1893	2109	G3.
FT	DISULFID	51	129	BY SIMILARITY.
FT	DISULFID	171	242	BY SIMILARITY.
FT	DISULFID	195	216	BY SIMILARITY.
FT	DISULFID	269	345	BY SIMILARITY.
FT	DISULFID	293	314	BY SIMILARITY.
FT	DISULFID	542	613	BY SIMILARITY.
FT	DISULFID	566	587	BY SIMILARITY.
FT	DISULFID	640	715	BY SIMILARITY.
FT	DISULFID	664	685	BY SIMILARITY.
FT	DISULFID	1859	1870	BY SIMILARITY.
FT	DISULFID	1864	1879	BY SIMILARITY.
FT	DISULFID	1881	1890	BY SIMILARITY.
FT	DISULFID	1897	1908	BY SIMILARITY.
FT	DISULFID	1925	2017	BY SIMILARITY.
FT	DISULFID	1993	2009	BY SIMILARITY.
FT	DISULFID	2024	2067	BY SIMILARITY.
FT	DISULFID	2053	2080	BY SIMILARITY.
FT	CARBOHYD	76	76	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	122	122	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	330	330	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	388	388	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	439	439	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	644	644	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	700	700	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	765	765	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	CARBOHYD	801	801	N-LINKED (GLCNAC. .) (POTENTIAL).
FT	VARSPIC	1856	1892	Missing (in isoform 2).
FT	CONFLICT	362	362	/FTid=VSP 003073.
FT	CONFLICT	362	362	E -> D (IN REF. 3).

Query Match 36.2%; Score 47; DB 1; Length 2109;  
Best Local Similarity 47.6%; Pred. No. 1.4e+02;  
Matches 10; Conservative 3; Mismatches 8; Indels 0; Gaps 0;

Qy 4 LPAYVGLSPGEQYHRGVGV 24  
Db 957 LVEVVTAAFGQERKGSIGV 977

RESULT 10  
COBI\_MYCTU  
ID COBI\_MYCTU STANDARD; PRT; 508 AA.

AC Q10677;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Cohalamin biosynthesis protein COBI [includes: Precorrin-2 C20-methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2 methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.1.-)]  
DE COBI OR COBI OR RV2066 OR MT2126 OR MTCY49.05 OR MB2092.  
OS Mycobacterium tuberculosis, and  
OS Mycobacterium bovis  
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;

—





DR PRINTS; PR00415; ACONITASE.  
 DR PRODOM; PD000511; Aconitase N; 1.  
 DR TIGRFAMS; TIGR01341; aconitase 1; 1.  
 DR PROSITE; PS00450; ACONITASE\_1; 1.  
 DR PROSITE; PS01244; ACONITASE\_2; 1.  
 KW Lyase; Tricarboxylic acid cycle; Iron-sulfur; 4Fe-4S.  
 FT METAL 499 499 IRON-SULFUR (4FE-4S) (BY SIMILARITY).  
 FT METAL 565 565 IRON-SULFUR (4FE-4S) (BY SIMILARITY).  
 FT METAL 568 568 IRON-SULFUR (4FE-4S) (BY SIMILARITY).  
 SQ SEQUENCE 961 AA; 104025 MW; 689566D85E31F596 CRC64;  
 Query Match 34.6%; Score 45; DB 1; Length 961;  
 Best Local Similarity 48.0%; Pred. No. 1.2e+02;  
 Matches 12; Conservative 5; Mismatches 6; Indels 2; Gaps 2;  
 QY 2 AGLPAVVGL-SPGEQYHRCGVGL 25  
 Db 927 AEFDAVVRITPGEADYIRNG-GIL 950  
 RESULT 14  
 ID ILVD CHRVO STANDARD; PRT; 618 AA.  
 AC Q7NYJ7;  
 DT 15-MAR-2004 (Rel. 43, Created)  
 DT 15-MAR-2004 (Rel. 43, Last sequence update)  
 DT 15-MAR-2004 (Rel. 43, Last annotation update)  
 DE Dihydroxy-acid dehydratase [EC 4.2.1.9] (DAD).  
 GN ILVD OR CV1277.  
 OS Chromobacterium violaceum.  
 OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;  
 OC Neisseriaceae; Chromobacterium.  
 OX NCBI\_TaxID=536;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 12472 / DSM 30191;  
 RX MEDLINE=22882880; PubMed=14500782;  
 RA Vasconcelos A.T.R., de Almeida D.F., Hungria M., Guimaraes C.T.,  
 RA Antonio R.V., Almeida F.C., de Almeida L.G.P., de Almeida R.,  
 RA Alves-Gomes J.A., Andrade E.M., Azarpe J., de Araujo M.F.F.,  
 RA Astolfi-Filho S., Azevedo V., Baptista A.J., Batais L.A.M.,  
 RA Batista J.S., Belo A., van den Berg C., Bogo M., Bonatto S.,  
 RA Bordignon J., Brigidio M.M., Brito C.A., Brocchi M., Burity H.A.,  
 RA Camargo A.A., Cardoso D.D.P., Carneiro N.P., Carrao S.M.,  
 RA Carvalho C.M.B., Cascardo J.C.M., Cavada B.S., Chueire L.M.O.,  
 RA Creczynski-pasa T.B., Cunha-Junior N.C., Fagundes N., Falcao C.L.,  
 RA Fantinatti F., Farias I.P., Felipe M.S.S., Ferrari L.P., Ferro J.A.,  
 RA Ferro M.I.T., Franco G.R., Freitas N.S.A., Furian L.R.,  
 RA Garzinelli R.T., Gomes E.A., Goncalves P.R., Grangeiro T.B.,  
 RA Grattapaglia D., Grisard E.C., Hanna E.S., Jardim S.N., Laurino J.,  
 RA Leoi L.C.T., Lima L.F.A., Loureiro M.F., Lyra M.C.C.P.,  
 RA Madeira H.M.F., Manfio G.P., Maranhao A.O., Martins W.S.,  
 RA di Mauro S.M.Z., de Medeiros S.R.B., Meissner R.V., Moreira M.A.M.,  
 RA Nascimento F.F., Nicolas M.F., Oliveira J.G., Oliveira S.C.,  
 RA Paixao R.F.C. Parente J.A., Piedra F.O., Pena S.D.J., Pereira J.O.,  
 RA Pereira M., Pinto L.S.R.C., Pintos L.S., Porto J.I.R., Potrich D.P.,  
 RA Ramalho-Neto C.E., Reis A.M.M., Rigo L.U., Rondinelli E.,  
 RA Santos E.B.P., Santos F.R., Schneider M.P.C., Seuneh H.N.,  
 RA Silva A.M.R., da Silva A.L.C., Silva D.W., Silva R., Simoes I.C.,  
 RA Simon D., Soares C.M.A., Soares R.B.A., Souza E.M., Souza K.R.L.,  
 RA Souza R.C., Steffens M.B.R., Steindel M., Teixeira S.R., Urmenyi T.,  
 RA Vettore A., Wassen R., Zaha A., Simpson A.J.G.;  
 RT "The complete genome sequence of Chromobacterium violaceum reveals  
 RT remarkable and exploitable bacterial adaptability";  
 RL Proc. Natl. Acad. Sci. U.S.A. 100:11660-11665(2003).  
 CC -!- CATALYTIC ACTIVITY: 2,3-dihydroxy-3-methylbutanoate = 3-methyl-2-oxobutanate + H(2)O.  
 CC -!- COFACTOR: Binds 1 4Fe-4S cluster (Potential).  
 CC -!- PATHWAY: Valine and isoleucine biosynthesis; fourth step.  
 CC -!- SIMILARITY: Belongs to the ilvd / edd family.  
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 CC -----  
 DR EMBL; AF016914; AAQ58952.1; -.  
 DR HAMAP; MF 00012; -; 1.  
 DR PROSITE; PS00886; ILVD EDD 1; 1.  
 DR PROSITE; PS00887; ILVD EDD 2; 1.  
 KW Branched-chain amino acid Biosynthesis; Lyase; Iron; Iron-sulfur;  
 KW 4Fe-4S; Complete proteome.  
 FT METAL 122 122 IRON-SULFUR (4FE-4S) (POTENTIAL).  
 FT METAL 195 195 IRON-SULFUR (4FE-4S) (POTENTIAL).  
 SQ SEQUENCE 618 AA; 65582 MW; 3273399ABAI92A05 CRC64;  
 Query Match 34.2%; Score 44.5; DB 1; Length 618;  
 Best Local Similarity 30.0%; Pred. No. 89;  
 Matches 9; Conservative 7; Mismatches 7; Indels 7; Gaps 1;  
 QY 3 GLPAVVGLSPGEQYHRCGVGL 25  
 Db 307 GVPCLSKVAPATQKYMEDVHRAGGVIGIL 336  
 RESULT 15  
 ID M18A HUMAN STANDARD; PRT; 2054 AA.  
 AC Q92614; O8IXP8;  
 DT 10-OCT-2003 (Rel. 42, Created)  
 DT 10-OCT-2003 (Rel. 42, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Myosin XVIIIa (Myosin 18A) (Myosin containing PDZ domain).  
 GN MYO18A OR MYSPDZ OR KIAA0216.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 OX NCBI\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A. (ISOFORM 1).  
 RC MEDLINE=97191544; PubMed=9039502;  
 RX Nagase T., Seki N., Ishikawa K.-I., Ohira M., Kawabayasi Y.,  
 RA Ohara O., Tanaka A., Kotani H., Miyajima N., Nomura N.;  
 RT "Prediction of the coding sequences of unidentified human genes. VI.  
 RT The coding sequences of 80 new genes (KIAA0201-KIAA0280) deduced by  
 RT analysis of cDNA clones from cell line KG-1 and brain.";  
 RL DNA Res. 3:321-329(1996).  
 RN [2]  
 RP SEQUENCE FROM N.A. (ISOFORM 3).  
 RC TISSUE=Testis;  
 RX MEDLINE=22388257; PubMed=12477932;  
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,  
 RA Klausner R.D., Colling F.S., Wagner L., Shenmen C.M., Schuler G.D.,  
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,  
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,  
 RA Datschenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,  
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,  
 RA Brownstein M.J., Usdin T.B., Tothiyuki S., Carninci P., Prange C.,  
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,  
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,  
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,  
 RA Villalón D.K., Murray D.M., Sodergren E.J., Lu X., Gibbs R.A.,  
 RA Fahy J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,  
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,  
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,  
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,  
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,  
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;  
 RT "Generation and initial analysis of more than 15,000 full-length human  
 RT and mouse cDNA sequences";  
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).  
 RN [3]

RX ALTERNATIVE SPLICING (ISOFORMS 1 AND 2).  
 RA MEDLINE=22646225; PubMed=12761286;  
 RA Mori K., Furusawa T., Okubo T., Inoue T., Ikawa S., Yanai N.,  
 RA Mori K.J., Obinata M.;  
 RT "Genome structure and differential expression of two isoforms of a  
 RT novel PDZ-containing myosin (MysPDZ) (Mys18A).";  
 RL J. Biochem. 133:405-413 (2003).  
 CC -!- FUNCTION: May be involved in the maintenance of the stromal cell  
 CC architectures required for cell to cell contact (By similarity).  
 CC -!- SUBCELLULAR LOCATION: Isoform 1 co-localizes with the  
 CC endoplasmatic reticulum-Golgi complex; isoform 2, which lacks the  
 CC PDZ domain, is diffusely localized in the cytoplasm (By  
 CC similarity).  
 CC -!- ALTERNATIVE PRODUCTS:  
 CC Event=Alternative splicing; Named isoforms=3;  
 CC Name=1; Synonyms=Alpha;  
 CC IsoId=Q92614-1; Sequence=Displayed;  
 CC Name=2; Synonyms=Beta;  
 CC IsoId=Q92614-2; Sequence=VSP\_007869, VSP\_007870;  
 CC Name=3;  
 CC IsoId=Q92614-3; Sequence=VSP\_007871, VSP\_007872;  
 CC Note=No experimental confirmation available;  
 CC -!- SIMILARITY: Contains 1 IQ domain.  
 CC -!- SIMILARITY: Contains 1 myosin-like globular head domain.  
 CC -!- SIMILARITY: Contains 1 PDZ/DHR domain.  
 CC -!- CAUTION: The TIAF1 protein is coded in the 3'UTR region of MysPDZ.  
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 -----  
 DR EMBL; D86970; BAAL3206.2; ALT INIT.  
 DR EMBL; BC039612; AAH39612.1; -.  
 DR PIR; PT0271; PT0271.  
 DR GO; GO:0005653; C:perinuclear space; TAS.  
 DR GO; GO:0006916; P:anti-apoptosis; TAS.  
 DR InterPro; IPR000048; IQ\_region.  
 DR InterPro; IPR003345; M\_repeat.  
 DR InterPro; IPR001609; myosin\_head.  
 DR InterPro; IPR001478; PDZ.  
 DR Pfam; PF00612; IQ; 1.  
 DR Pfam; PF02370; M; 7.  
 DR Pfam; PF00063; myosin\_head; 1.  
 DR Pfam; PF00595; PDZ; 1.  
 DR PRINTS; PR00193; MYOSINHEAVY.  
 DR PRODOM; PD000355; myosin\_head; 1.  
 DR SMART; SM00015; IQ; 1.  
 DR SMART; SM00242; MYSC; 1.  
 DR SMART; SM00228; PDZ; 1.  
 DR PROSITE; PS50096; IQ; 1.  
 DR PROSITE; PS50106; PDZ; 1.  
 DR Myosin; ATP-binding; Coiled coil; Alternative splicing.  
 DR DOMAIN 220 311 PDZ.  
 DR DOMAIN 420 1186 MYOSIN HEAD-LIKE.  
 DR DOMAIN 1188 1217 IQ.  
 DR DOMAIN 1246 1971 COILED COIL (POTENTIAL).  
 DR NP\_BIND 498 505 ATP (POTENTIAL).  
 DR VARSPPLIC 1 331 Missing (in isoform 2).  
 FT FTID=VSP\_007869.  
 FT SD -> MR (in isoform 2).  
 FT FTID=VSP\_007870.  
 FT Missing (in isoform 3).  
 FT FTID=VSP\_007871.  
 FT Missing (in isoform 3).  
 FT FTID=VSP\_007872.  
 FT SEQUENCE 2054 AA; 233113 MW; 52BFA0AA273E18F7 CRC64;  
 Query Match 34.2%; Score 44.5; DB 1; Length 2054;  
 Best Local Similarity 38.5%; Pred. No. 3.1e+02;

Matches 10; Conservative 3; Mismatches 4; Indels 9; Gaps 1;  
 QY 7 VVGLSPGEQF-----YHGGVG 23  
 Db 651 VLGISPDEQKACWFLAAYHLGAAG 676

Search completed: May 7, 2004, 12:34:30  
 Job time: 11 secs

01-JUN-2003 (T

```

DT 01-JUN-2003 (TReMBLrel. 24, Last sequence update)
DE 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
GN BLO130.
OS Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiyama T.,
RA Sasanoto S., Watanabe A., Idesawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpo S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;
RT "Complete genomic sequence of nitrogen-fixing symbiotic bacterium
RT Bradyrhizobium japonicum USDA110.";
RL DNA Res. 9:189-197(2002).
DR EMBL; AF005935; BAC45395.1; -.
DR InterPro; IPR005493; Methyltransf_6.
DR Pfam; PF03737; Methyltransf_6; 1.
KW Complete proteome.
SQ SEQUENCE 242 AA; 25216 MW; 588F9C0B9396414B CRC64;

Query Match 43.1%; Score 56; DB 16; Length 242;
Best Local Similarity 57.1%; Pred. No. 5.5;
Matches 12; Conservative 1; Mismatches 4; Indels 4; Gaps 1;

QY 1 MAGLPVVGLSPGQYHRGG 21
Db :||| ||| |||||
63 LARLFGVGLKP----YHRGG 79

RESULT 3
Q87YM5 PRELIMINARY; PRT; 330 AA.
AC Q87YM5;
DT 01-JUN-2003 (TReMBLrel. 24, Created)
DT 01-JUN-2003 (TReMBLrel. 24, Last sequence update)
DE 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
DE Conserved hypothetical protein.
GN PSPT03770.
OS Pseudomonas syringae (pv. tomato).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=323;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DC3000;
RA Buehl R., Joardar V., Khouri H., Fedorova N., Tran B., Russell D.,
RA Berry K., Utterback T., Van Aken S., Feldblyum T., Gwinn M.,
RA Dodson R., Deboy R., Durkin A., Kolonay J., Madupu R., Daugherty S.,
RA Brinkac L., Beanan M., Haft D., Selengut J., Nelson W., Davidsen T.,
RA White O., Fraser C., Collier A.;
RT "Complete sequence of Pseudomonas syringae.";
RT Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE016869; AAO57239.1; -.
DR TIGR; PSPT03770; -.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 330 AA; 36369 MW; 99876752E292FA71 CRC64;

Query Match 41.5%; Score 54; DB 16; Length 330;
Best Local Similarity 43.5%; Pred. No. 15;
Matches 10; Conservative 4; Mismatches 9; Indels 0; Gaps 0;

QY 3 GLPAVVGLSPGQYHRGGVGL 25
Db :||| :||| :|||
308 GQPLINGLAPSEAVFPGGIGKL 330

RESULT 4
Q8PE55 PRELIMINARY; PRT; 150 AA.
AC Q8PE55;
DT 01-OCT-2002 (TReMBLrel. 22, Created)
DT 01-OCT-2002 (TReMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)
DE Deoxycytidylate deaminase.
GN XCC0126.
OS Xanthomonas campestris (pv. campestris).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=340;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 33913 / NCPPB 528;
RX MEDLINE=2202145; PubMed=12024217;
RA da Silva A.C.R., Ferro J.A., Reinach F.C., Parah C.S., Furlan L.R.,
RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
RA Camarotte G., Cannavan F., Cardoso J., Chambergo F., Ciapina L.P.,
RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.,
RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
RA Setubal J.C., Kitajima J.P.;
RT "Comparison of the genomes of two Xanthomonas pathogens with differing
RT host specificities.";
RL Nature 417:459-463(2002).
DR EMBL; AE012107; AAM39445.1; -.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR InterPro; IPR002125; dCMP/cyt deam.
DR Pfam; PF00383; dCMP_cyt_deam; 1.
KW Complete proteome.
SQ SEQUENCE 150 AA; 16201 MW; 63D9C17D44DC8B43 CRC64;

Query Match 40.0%; Score 52; DB 16; Length 150;
Best Local Similarity 58.8%; Pred. No. 12;
Matches 10; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 2 AGLPAVVGLSPGQYEH 18
Db :||| :||| :|||
104 AGIKRVVALPGESEH 120

RESULT 5
Q8YBRO PRELIMINARY; PRT; 426 AA.
AC Q8YBRO;
DT 01-NOV-1999 (TReMBLrel. 12, Created)
DT 01-NOV-1999 (TReMBLrel. 12, Last sequence update)
DT 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
DE Hypothetical protein APE1539.
GN APE1539.
OS Aeropyrum pernix.
OC Archaea; Crenarchaeota; Thermoprotei; Desulfurococcales;
OC Desulfurococaceae; Aeropyrum.
OX NCBI_TaxID=56636;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K1;
RX MEDLINE=99310339; PubMed=10382966;
RA Kawarabayasi Y., Hino Y., Horikawa H., Yamazaki S., Haikawa Y.,
RA Jin-no K., Takahashi M., Sekine M., Baba S.-I., Ankai A., Kosugi H.,
RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudoh Y.,
RA Yamazaki J., Kishida N., Oguchi A., Aoki K.-I., Kubota K.,
RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.;

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RESULT 11	
Q9L525	PRELIMINARY; PRT; 758 AA.
AC	Q8L525;
ID	Q8L525;
DT	01-OCT-2002 (TrEMBLrel. 22, Created)
DT	01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT	01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DT	BL103C09.11 protein (P0451D05.19 protein).
GN	BL103C09.11 OR P0451D05.19.
OS	Oryza sativa (japonica cultivar-group).
OC	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC	Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC	Ehrhartoideae; Oryzeae; Oryza.
OX	NCBI_TaxID=39947;
OX	[1]
RP	SEQUENCE FROM N.A.
RP	STRAIN=cv. Nipponbare;
RC	STRAIN=cv. Nipponbare;
RA	Sasaki T., Matsumoto T., Yamamoto K.;
RA	"Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC
RT	clone:BL103C09."
RT	Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
RL	[2]
RN	SEQUENCE FROM N.A.
RP	STRAIN=cv. Nipponbare;
RC	STRAIN=cv. Nipponbare;
RA	Sasaki T., Matsumoto T., Yamamoto K.;
RA	"Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, PAC
RT	clone:P0451D05."
RT	Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
RL	EMBL; AF003333; BAB91821.1; -
DR	EMBL; AF003253; BAB92317.1; -
DR	Gramene; Q8L525; -
DR	SEQUENCE 758 AA; 81761 MW; 2A95E763198FE12C CRC64;
QY	Query Match 38.8%; Score 50.5; DB 10; Length 758;
DB	Best Local Similarity 32.4%; Pred. No. 1.2e+02;
	Matches 12; Conservative 3; Mismatches 7; Indels 15; Gaps
QY	3 GLPVGVLSPGEQBY-----HRGGVG 24
DB	552 GAAGAIGLAQEQNFGPTFALLPVMQFGSQHPGGVG 588
RESULT 12	
Q9KQK6	PRELIMINARY; PRT; 277 AA.
AC	Q9KQK6;
ID	Q9KQK6;
DT	01-OCT-2000 (TrEMBLrel. 15, Created)
DT	01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT	01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DT	Formyltetrahydrofolate deformylase.
GN	VC1992.
OS	Vibrio cholerae.
OC	Bacteria; Proteobacteria; Gammaproteobacteria; Vibrionales;
OC	Vibrionaceae; Vibrio.
OX	NCBI_TaxID=666;
OX	[1]
RN	SEQUENCE FROM N.A.
RP	STRAIN=El Tor N16961 / Serotype O1;
RC	MEDLINE=20406833; PubMed=10952301;
RA	Heidelberg J.F., Eisen J.A., Nelson W.C., Clayton R.A., Gwinn M.L.,
RA	Hodson R.J., Haft D.H., Hickey E.K., Peterson J.D., Umayam L.A.,
RA	Gill S.R., Nelson K.E., Read T.D., Tettelin H., Richardson D.,
RA	Ermolaeva M.D., Vamathevan J., Bass S., Qin H., Dragoi I., Sellers P.
RA	McDonald L., Utterback T., Fleischmann R.D., Nierman W.C., White O.,
RA	Salzberg S.L., Smith H.O., Colwell R.R., Mekalanos J.J., Venter J.C.,
RA	Fraser C.M.;
RA	"DNA sequence of both chromosomes of the cholera pathogen Vibrio
RT	cholerae.";
RL	Nature 406:477-483(2000).
DR	EMBL; AE004274; AAF95140.1; -
DR	PIR; F82130; F82130.
DR	HSSP; P08179; 2GAR.

```

DR TIGR; VC1992; -.
DR GO; GO:0016537; F:amino acid binding; IEA.
DR GO; GO:0008864; F:formyltetrahydrofolate deformylase activity; IEA.
DR GO; GO:0016742; F:hydroxymethyl-, formyl- and related transfe. . .; IEA.
DR GO; GO:0006189; P:de novo IMP biosynthesis; IEA.
DR GO; GO:0009058; P:biosynthesis; IEA.
DR InterPro; IPR002912; ACT.
DR InterPro; IPR002376; formyl_transf.
DR PFAM; PF01842; ACT; 1.
DR PFAM; PF00551; formyl_transf; 1.
DR PRINTS; PR01575; FFH4HYDRLASE.
DR TIGRFAMs; TIGR00655; PurU; 1.
DR Complete proteome.
SQ SEQUENCE 277 AA; 31373 MW; A703491654753DC6 CRC64;

Query Match 38.5%; Score 50; DB 16; Length 277;
Best Local Similarity 52.9%; Pred.No. 47;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

Qy 4 LPVVGLSPGGEQYHRG 20
||| :| :| :| :|
Db 190 LPAFIGAKPYQAYERG 206

RESULT 13
Q7VF15 PRELIMINARY; PRT; 284 AA.
AC Q7VF15;
DT 01-OCT-2003 (TRENBLrel. 25, Created)
DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Formyltetrahydrofolate deformylase PurU (EC 3.5.1.10).
GN PURU OR HH1691.
OS Helicobacter hepaticus.
OC Bacteria; Proteobacteria; Epsilonproteobacteria; Campylobacterales;
OC Helicobacteraceae; Helicobacter.
OX NCBI_TaxID=32025;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 51449 / 3B1;
RX MEDLINE=22709201; PubMed=12810954;
RA Suerbaum S., Josenhans C., Sterzenbach T., Drescher B., Brandt P.,
Bell M., Droege M., Fartmann B., Fischer H.-P., Ge Z., Hoerster A.,
Holland R., Klein K., Koenig J., Macko L., Mendz G.L., Nyakatura G.,
Schauer D.B., Shen Z., Weber J., Froesch M., Fox J.G.;
RA "The complete genome sequence of the carcinogenic bacterium
RT Helicobacter hepaticus."
RL Proc. Natl. Acad. Sci. U.S.A. 100:7901-7906(2003).
RW EMBL; AB017149; AAP78288.1; -.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 284 AA; 32660 MW; AAED2DC5086236E3 CRC64;

Query Match 38.5%; Score 50; DB 16; Length 284;
Best Local Similarity 52.9%; Pred.No. 49;
Matches 9; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

Qy 4 LPVVGLSPGGEQYHRG 20
||| :| :| :| :|
Db 197 LPAFIGANPYQAYERG 213

RESULT 14
Q82RR7 PRELIMINARY; PRT; 335 AA.
AC Q82RR7;
DT 01-JUN-2003 (TRENBLrel. 24, Created)
DT 01-JUN-2003 (TRENBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Putative terpene cyclase.
GN TPCL1 OR SAV76.
OS Streptomyces avermitilis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;

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OC Streptomycineae; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=33903;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=21477403; PubMed=11572948;
RA Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
Shinose M., Takahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
RA "Genome sequence of an industrial microorganism Streptomycetes
RT avermitilis: deducing the ability of producing secondary
RT metabolites."
RL Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=22608306; PubMed=12692562;
RA Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
Sakaki Y., Hattori M., Omura S.;
RA "Complete genome sequence and comparative analysis of the industrial
RT microorganism Streptomycetes avermitilis."
RL Nat. Biotechnol. 21:526-531(2003).
DR EMBL; AP005021; BAC67785.1; -.
DR InterPro; IPR008949; Terpenoid_synth.
DR Complete proteome.
RW SEQUENCE 335 AA; 36480 MW; 49B8477E2D52666F CRC64;

Query Match 38.5%; Score 50; DB 16; Length 335;
Best Local Similarity 47.6%; Pred.No. 58;
Matches 10; Conservative 3; Mismatches 8; Indels 0; Gaps 0;

Qy 3 GLPAVGLSPGGEQYHRGGVG 23
||| :| :| :| :|
Db 8 GLPAPAGISPGLEATRRHNLG 28

RESULT 15
Q9RR15 PRELIMINARY; PRT; 381 AA.
AC Q9RR15;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)
DE Cytochrome P450.
GN DR2473.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=RL / ATCC 13939 / DSM 20539 / NCIB 9279;
RX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
Vanatavan J.J., Lam P., McDonald L., Utterback T., Zaleski C.,
Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
Fraser C.M.;
RA "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans RL."
RL Science 286:1571-1577(1999).
CC -!- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.
DR EMBL; AE002076; AAF12016.1; -.
DR PIR; F75270; F75270.
DR TIGR; DR2473; -.
DR GO; GO:0004497; F:monooxygenase activity; IEA.
DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR001128; Cytochrome_P450.
DR Pfam; PF00067; P450; 1.
DR PRINTS; PR00385; P450.
DR PROSITE; PS00086; CYTOCHROME_P450; 1.

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KW Heme; Monooxygenase; Oxidoreductase; Complete proteome.  
 SQ SEQUENCE 381 AA; 41940 MW; F191EA69F1797B53 CRC64;

Query Match 38.5%; Score 50; DB 16; Length 381;  
 Best Local Similarity 100.0%; Pred. No. 67;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GLPAVVGLSP 12  
 |||||  
 Db 51 GLPAVVGLSP 60

Search completed: May 7, 2004, 12:37:50  
 Job time : 48.9167 secs



GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 69.75 seconds  
(without alignments)  
101.272 Million cell updates/sec

Title: US-09-786-214A-5  
Perfect score: 130  
Sequence: 1 MAGLPVAVGLSPGEQVHRGGVGL 25

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : A\_Geneseqp\_29Jan04.\*  
1: Geneseqp1980s.\*  
2: Geneseqp1990s.\*  
3: Geneseqp2000s.\*  
4: Geneseqp2001s.\*  
5: Geneseqp2002s.\*  
6: Geneseqp2003as.\*  
7: Geneseqp2003bs.\*  
8: Geneseqp2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	130	100.0	25	3 AAY84264	Aay84264 Peptide c
2	106	81.5	20	3 AAY84265	Aay84265 Truncated
3	75	57.7	15	3 AAY84269	Aay84269 Peptide d
4	72	55.4	14	3 AAY84266	Aay84266 Peptide d
5	68	52.3	13	3 AAY84267	Aay84267 Peptide d
6	65	50.0	13	3 AAY84268	Aay84268 Peptide d
7	53	40.8	394	4 AAU43816	AAU43816 Propionib
8	53	40.8	394	6 ABM40335	ABM40335 Propionib
9	52	40.0	187	6 ABU21296	ABU21296 Protein e
10	51.5	39.6	191	3 AAB69296	Aab69296 HIV-1 non
11	50	38.5	277	6 ABU49441	Abu49441 Protein e
12	49.5	38.1	192	3 AAB69298	Aab69298 HIV-1 non
13	49	37.7	278	6 ABU30626	Abu30626 Protein e
14	48.5	37.3	192	3 AAB69290	Aab69290 HIV-1 non
15	48	36.9	278	6 ABU39195	Abu39195 Protein e
16	48	36.9	287	6 ABU17217	Abu17217 Protein e
17	48	36.9	303	6 ADA36829	Ada36829 Acinetoba
18	47.5	36.5	114	4 AAE03295	Aae03295 Human gen
19	47.5	36.5	114	5 ABG64478	Abg64478 Human alb
20	47	36.2	151	5 ABU50765	Abu50765 Helicobac
21	47	36.2	155	5 ABU51750	Abu51750 Helicobac
22	47	36.2	238	6 ABU27483	Abu27483 Protein e
23	47	36.2	278	6 ABU44771	Abu44771 Protein e
24	47	36.2	280	6 ABU48054	Abu48054 Protein e
25	47	36.2	280	6 ABU31449	Abu31449 Protein e

26	47	36.2	280	6 ABU15054	Abu15054 Protein e
27	47	36.2	280	6 ABU47098	Abu47098 Protein e
28	47	36.2	282	6 ABU41187	Abu41187 Protein e
29	47	36.2	283	6 ABM69037	Abm69037 Photorhab
30	47	36.2	293	2 AAW98485	Aaw98485 H. pylori
31	47	36.2	293	6 ABU31073	Abu31073 Protein e
32	46.5	35.8	656	6 ABU27276	Abu27276 Protein e
33	46	35.4	46	3 AAY58745	Aay58745 Somatosta
34	46	35.4	143	5 AAO21677	Aao21677 Human sec
35	46	35.4	143	7 ADB64633	Adb64633 Human pro
36	46	35.4	184	5 ABP26551	Abp26551 Streptoco
37	46	35.4	234	4 AAB36208	Aab36208 Human imm
38	46	35.4	274	6 ABU26498	Abu26498 Protein e
39	46	35.4	287	7 ADB80057	Adb80057 Mycobacte
40	46	35.4	300	2 AAY16108	Aay16108 A formate
41	46	35.4	418	5 ABB04480	Abb04480 Arthrobac
42	46	35.4	450	5 ABB04479	Abb04479 Arthrobac
43	46	35.4	692	6 ABR40877	Abr40877 Oryza sat
44	46	35.4	921	2 AAY08304	Aay08304 Human col
45	46	35.4	1070	7 ADE08073	Ade08073 Novel pro

## ALIGNMENTS

RESULT 1  
AAY84264  
ID AAY84264 standard; peptide; 25 AA.  
XX  
AC AAY84264;  
XX  
DT 12-JUL-2000 (first entry)  
XX  
DE Peptide of alternate reading frame of macrophage colony stimulating gene.  
XX  
KW Renal cell carcinoma; antigen; cytotoxic T lymphocyte;  
KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
XX  
OS Homo sapiens.  
XX  
PN WO200013699-A1.  
XX  
PD 16-MAR-2000.  
XX  
PF 03-SEP-1999; 99WO-US020344.  
XX  
PA 04-SEP-1998; 98US-0099077P.  
(LUDW-) LUDWIG INST CANCER RES.  
PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
XX  
WPI; 2000-256859/22.  
N-PSDB; AAZ99672.  
Isolated polypeptide used to treat subjects having a disorder characterized by expression of alternative open reading frame macrophage-colony stimulating factor comprises 25 amino acid residue sequence.

Claim 1; Page 64; 74pp; English.

The present sequence represents a tumour rejection antigen precursor, and is encoded by an alternative open reading frame (ORF) of human macrophage colony stimulating gene. Peptides derived from the alternative ORF of macrophage-colony stimulating factor, when presented by an antigen presenting cell having a human leukocyte antigen (HLA) class I molecule, effectively induce the activation and proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF of macrophage-colony stimulating factor are useful for enriching selectively a population of T lymphocytes with CD8+ T lymphocytes. They are also useful for diagnosing a disorder characterized by expression of

CC the polypeptide, and for identifying functional variants and mimetics  
 XX  
 SQ Sequence 25 AA;

Query Match 100.0%; Score 130; DB 3; Length 25;  
 Best Local Similarity 100.0%; Pred. No. 2.3e-11;  
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAGLPAVVGSLSPGEQYHRGGVGL 25  
 |||||  
 Db 1 MAGLPAVVGSLSPGEQYHRGGVGL 25

RESULT 2  
 AAY84265  
 ID AAY84265 standard; peptide; 20 AA.

XX  
 AC AAY84265;  
 XX  
 DT 12-JUL-2000 (first entry)  
 XX  
 DE Truncated macrophage colony stimulating factor tumour antigen.  
 XX  
 KW tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX  
 OS Homo sapiens.  
 XX  
 PN WO200013699-A1.  
 XX  
 PD 16-MAR-2000.  
 XX  
 PF 03-SEP-1999; 99WO-US020344.  
 XX  
 PR 04-SEP-1998; 98US-0099077P.  
 XX  
 PA (LUDW-) LUDWIG INST CANCER RES.

XX  
 PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 XX  
 DR WPI; 2000-256859/22.  
 DR N-PSDB; AAZ99675.  
 XX  
 PT Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Claim 3; Page 64; 74pp; English.

XX  
 CC The present sequence represents a truncated tumour rejection antigen  
 CC precursor, and is encoded by a truncated alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX  
 SQ Sequence 20 AA;

Query Match 81.5%; Score 106; DB 3; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 4.7e-08;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAGLPAVVGSLSPGEQYHRG 20  
 |||||  
 Db 1 MAGLPAVVGSLSPGEQYHRG 20

RESULT 3  
 AAY84269

ID AAY84269 standard; peptide; 15 AA.

XX  
 AC AAY84269;  
 XX  
 DT 12-JUL-2000 (first entry)

XX  
 DE Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX  
 KW tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 PN WO200013699-A1.

XX  
 PD 16-MAR-2000.  
 XX  
 PF 03-SEP-1999; 99WO-US020344.  
 XX  
 PR 04-SEP-1998; 98US-0099077P.  
 XX  
 PA (LUDW-) LUDWIG INST CANCER RES.

XX  
 PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 XX  
 DR WPI; 2000-256859/22.

XX  
 PT Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX  
 CC The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX  
 SQ Sequence 15 AA;

Query Match 57.7%; Score 75; DB 3; Length 15;  
 Best Local Similarity 100.0%; Pred. No. 0.00087;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AGLPAVVGSLSPGEQ 16  
 |||||  
 Db 1 AGLPAVVGSLSPGEQ 15

RESULT 4  
 AAY84266

ID AAY84266 standard; peptide; 14 AA.

XX  
 AC AAY84266;  
 XX  
 DT 12-JUL-2000 (first entry)

XX  
 DE Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX  
 KW tumour rejection antigen; macrophage colony stimulating gene;

KW macrophage-colony stimulating factor; antigen presenting cell;  
XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Synthetic.  
OS Homo sapiens.  
XX WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 2; Page 39; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 14 AA;

Query Match 55.4%; Score 72; DB 3; Length 14;  
Best Local Similarity 100.0%; Pred. No. 0.0022;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 LPVVVGLSPGQEQY 17  
Db 1 LPVVVGLSPGQEQY 14

RESULT 5  
AAY84267  
ID AAY84267 standard; peptide; 13 AA.

AC AAY84267;

XX 12-JUL-2000 (first entry)

DE Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.  
OS Homo sapiens.

XX WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX

PR 04-SEP-1998; 98US-0099077P.  
XX (LUDW-) LUDWIG INST CANCER RES.  
PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
PI WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 52.3%; Score 68; DB 3; Length 13;  
Best Local Similarity 100.0%; Pred. No. 0.0074;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 PAVVGLSPGQEQY 17  
Db 1 PAVVGLSPGQEQY 13

RESULT 6  
AAY84268

ID AAY84268 standard; peptide; 13 AA.

AC AAY84268;

XX 12-JUL-2000 (first entry)

DE Peptide derived from macrophage colony stimulating gene alternative ORF.  
XX tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.  
OS Homo sapiens.

XX WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 50.0%; Score 65; DB 3; Length 13;  
Best Local Similarity 100.0%; Pred. No. 0.02;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 LPVVGLSPGEQE 16  
DB 1 LPVVGLSPGEQE 13

RESULT 7

AAU43816  
ID AAU43816 standard; protein; 394 AA.

XX AC AAU43816;

DT 13-FEB-2002 (first entry)

DE Propionibacterium acnes immunogenic protein #4712.

XX SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;  
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;  
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
KW dermatological; osteopathic; neuroprotectant.

XX Propionibacterium acnes.

PN WO200181581-A2.

XX 01-NOV-2001.

XX 20-APR-2001; 2001WO-US012865.

XX 21-APR-2000; 2000US-0199047P.

XX 02-JUN-2000; 2000US-0208941P.

XX 07-JUL-2000; 2000US-0216747P.

XX (CORI-) CORIXA CORP.

XX Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;

PI L'Maisonneuve J, Zhang Y, Jen S, Carter D;

XX WPI; 2001-616774/71.

XX N-PSDB; AAS59521.

XX Propionibacterium acnes polypeptides and nucleic acids useful for  
PT vaccinating against and diagnosing infections, especially useful for  
PT treating acne vulgaris.

XX Claim 6; SEQ ID NO 5011; 1069pp; English.

XX Sequences AAU3105-AAU68017 represent Propionibacterium acnes immunogenic  
CC polypeptides. The proteins and their associated DNA sequences are used in  
CC the treatment, prevention and diagnosis of medical conditions caused by  
CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,  
CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.  
CC P. acnes is also involved in infections of bone, joints and the central

CC nervous system, however it is particularly involved in the inflammatory  
CC lesions associated with acne vulgaris. A method for detecting the  
CC presence or absence of P. acnes in a patient comprises contacting a  
CC sample with a binding agent that binds to the proteins of the invention  
CC and determining the amount of bound protein in the sample. The  
CC polypeptides may be used as antigens in the production of antibodies  
CC specific for P. acnes proteins. These antibodies can be used to  
CC downregulate expression and activity of P. acnes polypeptides and  
CC therefore treat P. acnes infections. The antibodies may also be used as  
CC diagnostic agents for determining P. acnes presence, for example, by  
CC enzyme linked immunosorbent assay (ELISA). Note: The sequence data for  
CC this patent did not form part of the printed specification, but was  
CC obtained in electronic format directly from WIPO at  
CC ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 394 AA;

Query Match 40.8%; Score 53; DB 4; Length 394;  
Best Local Similarity 45.0%; Pred. No. 39;  
Matches 9; Conservative 5; Mismatches 6; Indels 0; Gaps 0;

QY 5 PAVVGLSPGEQEHYHGGVGV 24  
DB 255 PVVLGTAPGQGHDRHGTGV 274

RESULT 8

ABM40335

ID ABM40335 standard; protein; 394 AA.

XX AC ABM40335;

XX 20-OCT-2003 (first entry)

DE Propionibacterium acnes immunogenic polypeptide #5011.

XX Acne vulgaris; antiseborrheic; dermatological; antibacterial;  
KW immunostimulant; immune response; vaccine; immunogenic.

XX Propionibacterium acnes.

XX WO2003033515-A1.

XX 24-APR-2003.

XX 11-OCT-2002; 2002WO-US032727.

XX 15-OCT-2001; 2001US-00978825.

XX (CORI-) CORIXA CORP.

XX Mitcham JL, Skeiky YAW, Persing DH, Bhatia A, Maisonneuve JL;  
PI Zhang Y, Wang S, Jen S, Lodes MJ, Benson DR, Jones R, Carter D;

PI Barth B, Vallieve-Douglass J;

XX WPI; 2003-381789/36.

XX N-PSDB; ACF64450.

XX New Propionibacterium acnes polypeptides and polynucleotides encoding the  
PT polypeptide, useful for diagnosing, preventing or treating acne vulgaris,  
PT or for stimulating an immune response specific for a P. acnes protein.

XX Claim 6; SEQ ID NO 5011; 1481pp; English.

XX The invention relates to an isolated polynucleotide (ACF64435-ACF64733)  
CC encoding a Propionibacterium acnes protein. The invention also relates to  
CC polypeptides encoded by the polynucleotides (ABM35624-ABM64536) and to  
CC immunogenic fragments of P. acnes polypeptides. The invention  
CC additionally encompasses expression vectors and host cells comprising a  
CC polynucleotide of the invention; antibodies against polypeptides of the  
CC polynucleotide of the invention; fusion proteins comprising a polypeptide of the invention; a  
CC method for stimulating an immune response specific for a P. acnes  
CC polypeptide and an isolated T cell population comprising T cells prepared

via this method; a vaccine composition (comprising P. acnes polypeptides, polynucleotides, antibodies, fusion proteins, T cell populations, or antigen-presenting cells that express the polypeptide); a method and kit for detecting or determining the presence or absence of P. acnes in a patient; and a method for inhibiting the development of P. acnes in a patient. The P. acnes polypeptides, polynucleotides, antibodies, fusion proteins, T cell populations or antigen-presenting cells that express the polypeptides are useful for diagnosing, preventing or treating acne vulgaris, or for stimulating an immune response specific for a P. acnes protein. The polynucleotides can also be used as probes or primers for nucleic acid hybridisation. The vaccine composition is useful for the stimulation of an immune response against P. acnes, or for treating acne, and the kit is useful for performing a diagnostic assay. The present sequence represents a specifically claimed P. acnes polypeptide which is thought to contain an immunogenic region. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

Query Match 40.8%; Score 53; DB 6; Length 394;  
Best Local Similarity 45.0%; Pred. No. 39;  
Matches 9; Conservative 5; Mismatches 6; Indels 0; Gaps 0;

Qy 5 PAVVGLSPGEOYHRGGVGV 24  
Db 255 PVLGTAPGGQHRHGGV 274

RESULT 9  
ABU21296  
ID ABU21296 standard; protein; 187 AA.  
XX ABU21296;  
AC ABU21296;  
DT 19-JUN-2003 (first entry)  
DE Protein encoded by Prokaryotic essential gene #6823.  
KW Antisense; prokaryotic essential gene; cell proliferation; drug design.  
OS Burkholderia fungorum.  
PN WO200277183-A2.  
XX 03-OCT-2002.  
XX 21-MAR-2002; 2002WO-US009107.  
PR 21-MAR-2001; 2001US-00815242.  
PR 06-SEP-2001; 2001US-00948993.  
PR 25-OCT-2001; 2001US-0342923P.  
PR 08-FEB-2002; 2002US-00072851.  
PR 06-MAR-2002; 2002US-0362699P.  
XX (ELIT-) ELITRA PHARM INC.  
XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;  
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;  
XX WPI; 2003-029926/02.  
DR N-PSDB; ACA25166.  
XX New antisense nucleic acids, useful for identifying proteins or screening  
PT for homologous nucleic acids required for cellular proliferation to  
PT isolate candidate molecules for rational drug discovery programs.  
XX Claim 25; SEQ ID NO 49220; 1766pp; English.  
XX The invention relates to an isolated nucleic acid comprising any one of  
CC the 6213 antisense sequences given in the specification where expression  
CC of the nucleic acid inhibits proliferation of a cell. Also included are:

(1) a vector comprising a promoter operably linked to the nucleic acid encoding a polypeptide whose expression is inhibited by the antisense nucleic acid; (2) a host cell containing the vector; (3) an isolated polypeptide or its fragment whose expression is inhibited by the antisense nucleic acid; (4) an antibody capable of specifically binding the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular proliferation or the activity of a gene in an operon required for proliferation; (7) identifying a compound that influences the activity of the gene product or that has an activity against a biological pathway required for proliferation, or that inhibits cellular proliferation; (8) identifying a gene required for cellular proliferation or the biological pathway in which a proliferation-required gene or its gene product lies or a gene on which the test compound that inhibits proliferation of an organism acts; (9) manufacturing an antibiotic; (10) profiling a compound's activity; (11) a culture comprising strains in which the gene product is overexpressed or underexpressed; (12) determining the extent to which each of the strains is present in a culture or collection of strains; or (13) identifying the target of a compound that inhibits the proliferation of an organism. The antisense nucleic acids are useful for identifying proteins or screening for homologous nucleic acids required for cellular proliferation to isolate candidate molecules for rational drug discovery programs, or for screening homologous nucleic acids required for proliferation in cells other than S. aureus, S. typhimurium, K. pneumoniae or P. aeruginosa. The present sequence is encoded by one of the target prokaryotic essential genes. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

Query Match 40.0%; Score 52; DB 6; Length 187;  
Best Local Similarity 47.8%; Pred. No. 24;  
Matches 11; Conservative 2; Mismatches 6; Indels 4; Gaps 1;

Qy 3 GLPA-----VVLSPGEOYHRGG 21  
Db 48 GLPVLVNVVGMFGRTEQHAG 70

RESULT 10  
AAB69296  
ID AAB69296 standard; protein; 191 AA.  
XX AAB69296;  
XX 12-SEP-2003 (revised)  
DT 20-APR-2001 (first entry)  
DE HIV-1 non-subtype B clone 94CY032-3 vif protein.  
XX HIV-1; human immunodeficiency virus; non-subtype B; gag; pol; env; vpr; vif; vpr; tat; rev; nef; vaccine.  
XX Human immunodeficiency virus 1.  
XX WO200026416-A1.  
XX 11-MAY-2000.  
XX 25-OCT-1999; 99WO-US024837.  
XX 02-NOV-1998; 98US-00184418.  
XX (UABR-) UAB RES FOUND.  
XX Hahn BH, Shaw GM, Gao F;  
XX WPI; 2000-365651/31.  
XX Novel genomic nucleic acids of non-subtype B human immunodeficiency virus  
PT type 1 useful for detecting and treating AIDS comprises a specific  
PT nucleotide sequence.

XX Claim 41; Fig 16; 131pp; English.

XX The present in invention provides the protein and coding sequences for a

CC number of human immunodeficiency virus (HIV) type 1 non-subtype B

CC isolates. The sequences shown include the near full-length coding

CC sequences from each isolate, and the env, pol, vif, vpr, vpu, gag, tat,

CC rev and nef proteins. These can be used to detect the presence of HIV-1

CC in a sample and to produce antibodies against non-subtype B HIV-1 virus.

CC These antibodies can be used in vaccines to prevent and treat HIV

CC infection. (Updated on 12-SEP-2003 to standardise OS field)

XX Sequence 191 AA;

Query Match 39.6%; Score 51.5; DB 3; Length 191;

Best Local Similarity 58.8%; Pred. No. 29;

Matches 10; Conservative 3; Mismatches 3; Indels 1; Gaps 1;

QY 9 GLSPGEQYHRG-GVGV 24

|||:||||:||||:

Db 70 GLPGEQDHLGHGWSI 86

RESULT 11

ABU49441

ID ABU49441 standard; protein; 277 AA.

XX AC ABU49441;

XX 19-JUN-2003 (first entry)

XX Protein encoded by Prokaryotic essential gene #34968.

XX Antisense; prokaryotic essential gene; cell proliferation; drug design.

XX Vibrio cholerae.

OS WO200277183-A2.

PN 03-OCT-2002.

PD 21-MAR-2002; 2002WO-US009107.

PF 21-MAR-2001; 2001US-00815242.

PR 06-SEP-2001; 2001US-00948993.

PR 25-OCT-2001; 2001US-0342923P.

PR 08-FEB-2002; 2002US-00072851.

PR 06-MAR-2002; 2002US-0362699P.

XX (ELIT-) ELITRA PHARM INC.

PA Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;

PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX WPI; 2003-029926/02.

DR N-PSDB; ACA53311.

XX New antisense nucleic acids, useful for identifying proteins or screening

PT for homologous nucleic acids required for cellular proliferation to

PT isolate candidate molecules for rational drug discovery programs.

XX Claim 25; SEQ ID NO 77365; 1766pp; English.

XX The invention relates to an isolated nucleic acid comprising any one of

CC the 6213 antisense sequences given in the specification where expression

CC of the nucleic acid inhibits proliferation of a cell. Also included are:

CC (1) a vector comprising a promoter operably linked to the nucleic acid

CC encoding a polypeptide whose expression is inhibited by the antisense

CC nucleic acid; (2) a host cell containing the vector; (3) an isolated

CC polypeptide or its fragment whose expression is inhibited by the

CC antisense nucleic acid; (4) an antibody capable of specifically binding

CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular

CC proliferation or the activity of a gene in an operon required for

CC proliferation; (7) identifying a compound that influences the activity of

CC the gene product or that has an activity against a biological pathway

CC required for proliferation, or that inhibits cellular proliferation; (8)

CC identifying a gene required for cellular proliferation or the biological

CC pathway in which a proliferation-required gene or its gene product lies

CC or a gene on which the test compound that inhibits proliferation of an

CC organism acts; (9) manufacturing an antibiotic; (10) profiling a

CC compound's activity; (11) a culture comprising strains in which the gene

CC product is overexpressed or underexpressed; (12) determining the extent

CC to which each of the strains is present in a culture or collection of

CC strains; or (13) identifying the target of a compound that inhibits the

CC proliferation of an organism. The antisense nucleic acids are useful for

CC identifying proteins or screening for homologous nucleic acids required

CC for cellular proliferation to isolate candidate molecules for rational

CC drug discovery programs, or for screening homologous nucleic acids

CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,

CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is encoded by one of

CC the target prokaryotic essential genes. Note: The sequence data for this

CC patent did not form part of the printed specification, but was obtained

CC in electronic format directly from WIPO at

CC ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 277 AA;

Query Match 38.5%; Score 50; DB 6; Length 277;

Best Local Similarity 52.9%; Pred. No. 71;

Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPVVGLSPGEQYHRG 20

||||:||||:

Db 190 LPAFIGAKPYQQAYERG 206

RESULT 12

AAB69298

ID AAB69298 standard; protein; 192 AA.

XX AC AAB69298;

XX 12-SEP-2003 (revised)

DT 20-APR-2001 (first entry)

XX HIV-1 non-subtype B clone 96ZM751-3 vif protein.

XX HIV-1; human immunodeficiency virus; non-subtype B; gag; pol; env; vpu;

KW vif; vpr; tat; rev; nef; vaccine.

XX Human immunodeficiency virus 1.

OS WO200026416-A1.

XX 11-MAY-2000.

XX 25-OCT-1999; 99WO-US024837.

XX 02-NOV-1998; 98US-00184418.

XX (UABR-) UAB RES FOUND.

XX Hahn BH, Shaw GM, Gao F;

XX WPI; 2000-365651/31.

XX Novel genomic nucleic acids of non-subtype B human immunodeficiency virus

PT type 1 useful for detecting and treating AIDS comprises a specific

PT nucleotide sequence.

XX Claim 41; Fig 16; 131pp; English.

XX The present in invention provides the protein and coding sequences for a

CC number of human immunodeficiency virus (HIV) type 1 non-subtype B

CC isolates. The sequences shown include the near full-length coding

CC sequences from each isolate, and the env, pol, vif, vpr, vpu, gag, tat,

CC rev and nef proteins. These can be used to detect the presence of HIV-1  
CC in a sample and to produce antibodies against non-subtype B HIV-1 virus.  
CC These antibodies can be used in vaccines to prevent and treat HIV  
CC infection. (Updated on 12-SEP-2003 to standardise OS field)  
XX  
XX Sequence 192 AA;  
SQ

Query Match 38.1%; Score 49.5; DB 3; Length 192;  
Best Local Similarity 58.8%; Pred. NO. 56;  
Matches 10; Conservative 3; Mismatches 3; Indels 1; Gaps 1;

QY 9 GLSPGEGYHRG-GVGV 24  
|||:::||:  
Db 71 GLHPGEREWHLGHGVS I 87

RESULT 13  
ABU30626  
ID ABU30626 standard; protein; 278 AA.  
XX  
XX AC ABU30626;  
XX  
XX DT 19-JUN-2003 (first entry)  
XX  
XX DE Protein encoded by Prokaryotic essential gene #16153.  
XX  
XX KW Antisense; prokaryotic essential gene; cell proliferation; drug design.  
XX  
XX OS Haemophilus influenzae.  
XX  
XX PN W0200277183-A2.  
XX  
XX PD 03-OCT-2002.  
XX  
XX PF 21-MAR-2002; 2002WO-US009107.  
XX  
XX PR 21-MAR-2001; 2001US-00815242.  
XX PR 06-SEP-2001; 2001US-00948993.  
XX PR 25-OCT-2001; 2001US-0342923P.  
XX PR 08-FEB-2002; 2002US-00072851.  
XX PR 06-MAR-2002; 2002US-0362699P.  
XX  
XX PA (ELIT-) ELITRA PHARM INC.  
XX  
XX PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;  
XX PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;  
XX  
XX DR WPI; 2003-029926/02.  
XX DR N-PSDB; ACA34496.  
XX  
XX PT New antisense nucleic acids, useful for identifying proteins or screening  
XX PT for homologous nucleic acids required for cellular proliferation to  
XX PT isolate candidate molecules for rational drug discovery programs.

Claim 25; SEQ ID NO 58550; 1766pp; English.

The invention relates to an isolated nucleic acid comprising any one of  
the 6213 antisense sequences given in the specification where expression  
of the nucleic acid inhibits proliferation of a cell. Also included are:  
(1) a vector comprising a promoter operably linked to the nucleic acid  
encoding a polypeptide whose expression is inhibited by the antisense  
nucleic acid; (2) a host cell containing the vector; (3) an isolated  
polypeptide or its fragment whose expression is inhibited by the  
antisense nucleic acid; (4) an antibody capable of specifically binding  
the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular  
proliferation or the activity of a gene in an operon required for  
proliferation; (7) identifying a compound that influences the activity of  
the gene product or that has an activity against a biological pathway  
required for proliferation, or that inhibits cellular proliferation; (8)  
identifying a gene required for cellular proliferation or the biological  
pathway in which a proliferation-required gene or its gene product lies  
or a gene on which the test compound that inhibits proliferation of an  
organism acts; (9) manufacturing an antibiotic; (10) profiling a

```

CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than S. aureus, S. typhimurium,
CC K. pneumoniae or P. aeruginosa. The present sequence is encoded by one of
CC the target prokaryotic essential genes. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
XX
SQ Sequence 278 AA;
Query Match 36.9%; Score 48; DB 6; Length 278;
Best Local Similarity 52.9%; Pred. No. 1.4e+02;
Matches 9; Conservative 1; Mismatches 7; Indels 0; Gaps 0;
QY 4 LPAWVGLSPGEQYHRG 20
||| : ||| |||
Db 191 LPAFIGAKPYHQAYERG 207

Search completed: May 7, 2004, 12:33:42
Job time : 71.75 secs

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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 7.2 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-9

Perfect score: 106

Sequence: 1 MAGLPAVVGSLSPGQEXHRG 20

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 segs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	49	46.2	278	1 PURU_HAEIN	Q03432 haemophilus
2	48	45.3	780	1 PPSA_DEIRA	O83026 deinotholob
3	47	44.3	280	1 PURU_ECOL6	P38480 escherichia
4	47	44.3	280	1 PURU_ECOL1	P37051 escherichia
5	46	43.4	508	1 COBI_MYCTU	Q10677 mycobacteri
6	44	41.5	1172	1 CNA2_MOUSE	Q80xc6 mus musculu
7	44	41.5	3664	1 MINT_HUMAN	Q98t58 homo sapien
8	43	40.6	211	1 COBL_METUA	Q58917 methanococc
9	43	40.6	326	1 MER_METTI	Q9uxp0 methanolobu
10	43	40.6	390	1 COBL_MYCTU	Q10671 mycobacteri
11	43	40.6	505	1 Y76J_CABEL	P90838 caenorhabdi
12	43	40.6	516	1 C4AD_DROME	P94t33 drosophila
13	43	40.6	544	1 CH60_PROAC	Q9k2u4 propionibac
14	43	40.6	628	1 HNFA_MOUSE	P22361 mus musculu
15	43	40.6	628	1 HNFA_RAT	P15257 rattus norv
16	43	40.6	631	1 HNFA_HUMAN	P20823 homo sapien
17	43	40.6	633	1 PLB5_SCHPO	Q9v7n6 schizosach
18	43	40.6	658	1 VGI8_BPT4	P13332 bacterioph
19	43	40.6	677	1 SGI_HUMAN	P05060 homo sapien
20	43	40.6	983	1 RPOB_SUNNV	Q9qaz8 striped jac
21	42	39.6	400	1 TRUD_METKA	O8txj7 methanopyru
22	42	39.6	469	1 GLYC_YEAST	P37291 saccharomyc
23	42	39.6	808	1 PLB8_VIBPA	O87kn0 vibrio para
24	42	39.6	876	1 TOP1_VIBCH	O9krb2 vibrio chol
25	42	39.6	1065	1 RPOB_WARPO	P05272 marchantia
26	42	39.6	1712	1 CA24_HUMAN	P08572 homo sapien
27	41	38.7	156	1 RUVX_CAUCR	Q9ask8 caulobacter
28	41	38.7	213	1 MDCG_XANAC	O8ppw9 xanthomonas
29	41	38.7	243	1 SUMT_SNP7	P43451 synechococc
30	41	38.7	479	1 PTSE_VIBAL	P22825 vibrio algi
31	41	38.7	481	1 GYRB_FLEJA	Q9fax2 flexibacter
32	41	38.7	520	1 AT15_YEAST	P25641 saccharomyc
33	41	38.7	579	1 NH71_CABEL	Q9gtd4 caenorhabdi

34	41	38.7	774	1 LOL2_HUMAN	Q9v4k0 homo sapien
35	41	38.7	934	1 OD01_COXBU	P51056 coxiella bu
36	41	38.7	981	1 RRPO_AHNNV	Q9aic5 atlantic ha
37	40.5	38.2	387	1 OMA2_NEIMC	P18194 neisseria m
38	40.5	38.2	392	1 OMA2_NEIMC	Q51240 neisseria m
39	40.5	38.2	393	1 OMA1_NEIMC	P13415 neisseria m
40	40.5	38.2	395	1 OMA1_NEIMA	P57041 neisseria m
41	40.5	38.2	811	1 PRIA_RHOKU	P05445 rhodospiril
42	40	37.7	157	1 RISB_PYRFU	Q8u418 pyrococcus
43	40	37.7	193	1 VIF_SIVCZ	P17284 chimpanzee
44	40	37.7	316	1 Y273_SYNY3	P73894 synechocyst
45	40	37.7	331	1 KDD2_STRAW	Q82bx4 streptomyce

#### ALIGNMENTS

RESULT 1  
PURU\_HAEIN STANDARD; PRT; 278 AA.  
AC Q03432;  
DT 01-OCT-1994 (Rel. 30, Created)  
DT 01-NOV-1995 (Rel. 32, Last sequence update)  
DT 28-FEB-2003 (Rel. 41, Last annotation update)  
DE Formyltetrahydrofolate deformylase (EC 3.5.1.10) (Formyl-PH(4)  
DE hydrolase).  
GN PURU OR H11588.  
OS Haemophilus influenzae.  
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;  
OC Pasteurellaceae; Haemophilus.  
OX NCBI\_TaxID=727;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=Rd / KW20 / ATCC 51907;  
RX MEDLINE=95350630; PubMed=7542800;  
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,  
RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,  
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,  
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,  
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,  
RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,  
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,  
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,  
RA Venter J.C.;  
RT "Whole-genome random sequencing and assembly of Haemophilus influenzae  
RT Rd.";  
RL Science 269:496-512 (1995).  
RN [2]  
RP SEQUENCE OF 64-278 FROM N.A.  
RC STRAIN=RM 7004 / Serotype B;  
RX MEDLINE=93328119; PubMed=8335255;  
RA Maskell D.J.;  
RT "Cloning and sequencing of the Haemophilus influenzae *aroA* gene.";  
RL Gene 129:155-156 (1993).  
CC -!- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES  
THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF  
5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC  
GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL  
(BY SIMILARITY).  
CC -!- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate +  
tetrahydrofolate.  
CC -!- ENZYME REGULATION: Activated by methionine, inhibited by glycine  
(By similarity).  
CC -!- PATHWAY: De novo purine biosynthesis.  
CC -!- SUBUNIT: Homohexamer (By similarity).  
CC -!- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).  
-----  
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CC or send an email to license@sib-sib.ch).
CC -----
CC EMBL; U32833; AAC23236.1; -.
CC EMBL; L04686; AAA24942.1; -.
CC DR PIR; E64131; E64131.
CC DR HSSP; P08179; IGRC.
CC DR TIGR; H11588; -.
CC DR InterPro; IPR002912; ACT.
CC DR InterPro; IPR002376; formyl_transf.
CC DR InterPro; IPR004810; PurU.
CC DR Pfam; PF01842; ACT; 1.
CC DR Pfam; PF00551; formyl_transf; 1.
CC DR PRINTS; PR01575; FFF4HYDRLASE.
CC DR TIGRFAMs; TIGR00655; PurU; 1.
CC KW Purine biosynthesis; Hydrolase; One-carbon metabolism;
CC Complete proteome.
CC FT ACT_SITE 223 BY SIMILARITY.
CC FT CONFLICT 115 117 VIG -> RNR (IN REF. 2).
CC FT CONFLICT 138 140 HEN -> PK (IN REF. 2).
CC FT CONFLICT 205 205 K -> E (IN REF. 2).
CC SQ SEQUENCE 278 AA; 32173 MW; 7F375AB3C422EC4B CRC64;

Query Match 46.2%; Score 49; DB 1; Length 278;
Best Local Similarity 52.9%; Pred. No. 3.6;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPVVGLSPGCEQYVHRG 20
DB 191 LPFAFGAKPYQAYKRG 207

RESULT 2
PPSA DEIRA
ID _PPSA DEIRA STANDARD; PRT; 780 AA.
AC O83026.
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Phosphoenolpyruvate synthase (EC 2.7.9.2) (Pyruvate, water dikinase)
DE (PEP synthase).
DE PPSA OR DRI727.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
EX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Pauphilet W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Utterback T., Zalewski C.,
RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RA "Genome sequence of the radioresistant bacterium Deinococcus
RA radiodurans R1.";
RL Science 286:1571-1577(1999).
RN [2]
RP SEQUENCE OF 259-780 FROM N.A.
RA Narumi I., Islam S., Cherdchu K., Kikuchi M., Watanabe H.,
RA Kitayama S., Yamamoto K.;
RT "IS8101: the second insertion sequence element from Deinococcus
RT radiodurans.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -|- CATALYTIC ACTIVITY: ATP + pyruvate + H(2)O = AMP +
CC phosphoenolpyruvate + phosphate.
CC -|- PATHWAY: ESSENTIAL STEP IN GLUCOGENESIS WHEN PYRUVATE AND
CC LACTATE ARE USED AS A CARBON SOURCE.
CC -|- SIMILARITY: Belongs to the PEP-utilizing enzyme family.
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CC -----
CC DR EMBL; AE002014; AAF11283.1; -.
CC DR EMBL; AB016803; BAA32387.1; -.
CC DR PIR; D75361; D75361.
CC DR PIR; T44369; T44369.
CC DR TIGR; DRI727; -.
CC DR InterPro; IPR008279; PEP mobile.
CC DR InterPro; IPR006318; PEP_P trans.
CC DR InterPro; IPR006319; PEP synth.
CC DR InterPro; IPR000121; PEP utilizers.
CC DR InterPro; IPR002192; PPD_K N term.
CC DR Pfam; PF00391; PEP-utilizers; 1.
CC DR Pfam; PF02896; PEP-utilizers; 1.
CC DR Pfam; PF01326; PPD_K N term; 1.
CC DR PRINTS; PR01736; PHPTNFRASE.
CC DR PRODOM; PD000940; PEP utilizers; 1.
CC DR TIGRFAMs; TIGR01418; PEP synth; 1.
CC DR PROSITE; PS00370; PEP_ENZYMES_PHOS_SITE; 1.
CC DR PROSITE; PS00742; PEP_ENZYMES_2; 1.
CC KW Transferase; Kinase; ATP-binding; Phosphorylation; Complete proteome.
CC FT ACT_SITE 409 409 TELE-PHOSPHOHISTIDINE INTERMEDIATE
CC (BY SIMILARITY).
CC FT MOD_RES 409 409 PHOSPHORYLATION (BY SIMILARITY).
CC SQ SEQUENCE 780 AA; 84895 MW; AD555076324ADA47 CRC64;

Query Match 45.3%; Score 48; DB 1; Length 780;
Best Local Similarity 50.0%; Pred. No. 15;
Matches 9; Conservative 2; Mismatches 7; Indels 0; Gaps 0;

QY 3 GLPAVVGSLSPGCEQYVHRG 20
DB 418 GIPAVVGNGNATRELHNG 435

RESULT 3
PURU ECOL6
ID _PURU ECOL6 STANDARD; PRT; 280 AA.
AC P38480.
DT 01-OCT-1994 (Rel. 30, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Formyltetrahydrofolate deformylase (EC 3.5.1.10) (Formyl-FH(4)
DE hydrolase).
GN PURU OR C1696 OR SF1232 OR S1318.
OS Escherichia coli O6, and
OS Shigella flexneri.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=217992, 623;
RN [1]
RP SEQUENCE FROM N.A.
RC SPECIES=E.coli; STRAIN=O6:H1 / CFT073 / ATCC 700928;
RX MEDLINE=22388234; PubMed=12471157;
RA Welch R.A., Burland V., Plunkett G. III, Redford P., Roesch P.,
RA Rasko D., Buckles E.L., Liou S.-R., Boutin A., Hackett J., Stroud D.,
RA Mayhew G.F., Rose D.J., Zhou S., Schwartz D.C., Perna N.T.,
RA Mobley H.L.T., Donnenberg M.S., Blattner F.R.;
RT "Extensive mosaic structure revealed by the complete genome sequence
RT of uropathogenic Escherichia coli.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:17020-17024(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC SPECIES=S.flexneri; STRAIN=2457T / ATCC 700930 / Serotype 2a;
RX MEDLINE=93023838; PubMed=1406252;
RA Homococky A.E., Tucker S.C., Mauriello A.T.;
RA "Temperature regulation of Shigella virulence: identification of the
RA repressor gene virR, an analogue of hns, and partial complementation
RT -----
```

RT by tyrosyl transfer RNA (tRNA<sup>Tyr</sup>).";  
 RL Mol. Microbiol. 6:2113-2124(1992).  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RC SPECIES=S.flexneri; STRAIN=301 / Serotype 2a;  
 RX MEDLINE=22272406; PubMed=12384590;  
 RA Jin Q., Yuan Z., Xu J., Wang Y., Shen Y., Lu W., Wang J., Liu H.,  
 RA Yang J., Yang F., Zhang X., Zhang J., Yang G., Wu H., Qu D., Dong J.,  
 RA Sun L., Xue X., Zhao X., Gao Y., Zhu J., Kan B., Ding K., Chen S.,  
 RA Cheng H., Yao Z., He B., Chen R., Ma D., Qiang B., Wen Y., Hou Y.,  
 RA Yu J.;  
 RT "Genome sequence of Shigella flexneri 2a: insights into pathogenicity  
 RT through comparison with genomes of Escherichia coli K12 and O157.";  
 RL Nucleic Acids Res. 30:4432-4441(2002).  
 RN [4]  
 RP SEQUENCE FROM N.A.  
 RC SPECIES=S.flexneri; STRAIN=2457T / ATCC 700930 / Serotype 2a;  
 RX MEDLINE=22590274; PubMed=12704152;  
 RA Wei J., Goldberg M.B., Burland V., Venkatesan M.M., Deng W.,  
 RA Fournier G., Mayhew G.F., Plunkett G. III, Rose D.J., Darling A.,  
 RA Mau B., Perna N.T., Payne S.M., Runyen-Vance L.J., Zhou S.,  
 RA Schwartz D.C., Blattner F.R.;  
 RT "Complete genome sequence and comparative genomics of Shigella  
 RT flexneri serotype 2a strain 2457T.";  
 RL Infect. Immun. 71:2775-2786(2003).  
 RN [5]  
 RP IDENTIFICATION.  
 RX MEDLINE=94042872; PubMed=8226647;  
 RA Nagy P.L., McCorkle G., Zalkin H.;  
 RT "puru", a source of formate for purT-dependent phosphoribosyl-N-  
 RT formylglycinamide synthesis.";  
 RL J. Bacteriol. 175:7066-7073(1993).  
 CC -!- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES  
 CC THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF  
 CC 5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC  
 CC GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL.  
 CC -!- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate +  
 CC tetrahydrofolate.  
 CC -!- ENZYME REGULATION: Activated by methionine, inhibited by glycine  
 CC (by similarity).  
 CC -!- PATHWAY: De novo purine biosynthesis.  
 CC -!- SUBUNIT: Homohexamer (By similarity).  
 CC -!- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).  
 CC -----  
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 CC -----  
 DR EMBL; AE016760; AAN80163.1; -;  
 DR EMBL; X66849; -; NOT ANNOTATED\_CDS.  
 DR EMBL; AE015150; AAN42845.1; -;  
 DR EMBL; AE016982; AAP16730.1; -;  
 DR HSSP; P08179; 1GPC  
 DR InterPro; IPR002912; ACT.  
 DR InterPro; IPR002376; formyl\_transf.  
 DR InterPro; IPR004810; PurnU.  
 DR Pfam; PF01842; ACT; 1.  
 DR Pfam; PF00551; formyl\_transf; 1.  
 DR PRINTS; PR01575; FFH4HYDRLASE.  
 DR TIGRPMs; TIGR00655; PurnU; 1.  
 KW Purine biosynthesis; Hydrolyase; One-carbon metabolism;  
 KW Complete proteome.  
 FT ACT SIZE 225  
 FT CONFLICT 44 44 R -> L (IN REF. 2).  
 FT SEQUENCE 280 AA; 31920 MW; 55BC16B62727A419 CRC64;  
 SQ  
 Query Match 44.3%; Score 47; DB 1; Length 280;  
 Best Local Similarity 52.9%; Pred. No. 7.4;  
 Matches 9; Conservative 1; Mismatches 7; Indels 0; Gaps 0;

QY 4 LPVVVGLSPGEQYHRG 20  
 ||| : ||| |||  
 Db 193 LPAFIGARPYHQAYERG 209  
 ||| : ||| |||  
 RESULT 4  
 PURU\_ECOLI  
 ID PURU\_ECOLI STANDARD; PRT; 280 AA.  
 AC P37051;  
 DT 01-JUN-1994 (Rel. 29, Created)  
 DT 01-JUN-1994 (Rel. 29, Last sequence update)  
 DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 DE Formyltetrahydrofolate deformylase (EC 3.5.1.10) (Formyl-FH(4)  
 DE hydrolyase).  
 GN PURU OR TGS OR B1232.  
 OS Escherichia coli.  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;  
 OC Enterobacteriaceae; Escherichia.  
 OX NCBI\_TaxID=562;  
 RN [1]  
 RP SEQUENCE FROM N.A., AND SEQUENCE OF 1-6.  
 RC STRAIN=K12;  
 RX MEDLINE=94042872; PubMed=8226647;  
 RA Nagy P.L., McCorkle G., Zalkin H.;  
 RT "puru", a source of formate for purT-dependent phosphoribosyl-N-  
 RT formylglycinamide synthesis.";  
 RL J. Bacteriol. 175:7066-7073(1993).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=K12 / MG1655;  
 RX MEDLINE=94110230; PubMed=8282700;  
 RA Boesl M., Kersten H.;  
 RT "Organization and functions of genes in the upstream region of tyrt  
 RT of Escherichia coli: phenotypes of mutants with partial deletion of a  
 RT new gene [tgs].";  
 RL J. Bacteriol. 176:221-231(1994).  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=K12 / MG1655;  
 RX MEDLINE=97426617; PubMed=9278503;  
 RA Blattner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,  
 RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,  
 RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,  
 RA Mau B., Shao Y.;  
 RT "The complete genome sequence of Escherichia coli K-12.";  
 RL Science 277:1453-1474(1997).  
 RN [4]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=K12;  
 RX MEDLINE=97061202; PubMed=8905232;  
 RA Oshima T., Aiba H., Baba T., Fujita K., Hayashi K., Honjo A.,  
 RA Ikemoto K., Inada T., Itoh T., Kajihara M., Kanai K., Kashimoto K.,  
 RA Kimura S., Kitagawa M., Makino K., Masuda S., Miki T., Mizobuchi K.,  
 RA Mori H., Motomura K., Nakamura Y., Nishimoto H., Nishio Y., Saito N.,  
 RA Sampei G., Seki Y., Tagami H., Takemoto K., Wada C., Yamamoto Y.,  
 RA Yano M., Horiuchi T.;  
 RT "A 718-kb DNA sequence of the Escherichia coli K-12 genome  
 RT corresponding to the 12.7-28.0 min region on the linkage map.";  
 RL DNA Res. 3:137-155(1996).  
 RN [5]  
 RP CHARACTERIZATION.  
 RX MEDLINE=95173107; PubMed=7868604;  
 RA Nagy P.L., Marolewski A., Benkovic S.J., Zalkin H.;  
 RT "Formyltetrahydrofolate hydrolyase, a regulatory enzyme that functions  
 RT to balance pools of tetrahydrofolate and one-carbon tetrahydrofolate  
 RT adducts in Escherichia coli.";  
 RL J. Bacteriol. 177:1292-1298(1995).  
 CC -!- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES  
 CC THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF  
 CC 5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC  
 CC GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL.  
 CC -!- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate +

```

CC tetrahydrofolate.
CC -!- ENZYME REGULATION: Activated by methionine, inhibited by glycine.
CC -!- PATHWAY: De novo purine biosynthesis.
CC -!- SUBUNIT: Homohexamer.
CC -!- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).
CC -----
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CC -----
DR EMBL; L20251; AAC36846.1; -.
DR EMBL; M64675; AAA16860.1; ALT_INIT.
DR EMBL; AE000221; AAC74314.1; -.
DR EMBL; D90758; BAA36100.1; -.
DR EMBL; D90759; BAA36112.1; -.
DR EMBL; D90852; BAA16026.1; -.
DR PIR; C36871; C36871.
DR EcoGene; EGI1819; purU.
DR InterPro; IPR002912; ACT.
DR InterPro; IPR002376; formyl_transf.
DR InterPro; IPR004810; PurU.
DR Pfam; PF01842; ACT; 1.
DR Pfam; PF00551; formyl_transf; 1.
DR PRINTS; PR01575; FFH4HYDLRLASE.
DR TIGRFAMs; TIGR00655; PurU; 1.
KW Purine biosynthesis; Hydrolase; One-carbon metabolism;
KW Complete proteome.
FT ACT_SITE 225 BY SIMILARITY.
SQ SEQUENCE 280 AA; 31934 MW; 5667406D727A2C2 CRC64;

Query Match 44.3%; Score 47; DB 1; Length 280;
Best Local Similarity 52.9%; Pred. No. 7.4;
Matches 9; Conservative 1; Mismatches 7; Indels 0; Gaps 0;

QY 4 LPVVVGLSPGQEQYVRG 20
DB 193 LPFAFTGARGPHQAVERG 209

RESULT 5
ID COBI MYCTU STANDARD; PRT; 508 AA.
AC Q10677;
DT 15-JUL-1998 (Rel. 36, Created)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cobalamin biosynthesis protein COBIJ [Includes: Precorrin-2 C20-
DE methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2
DE methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.-)]].
GN COBIJ OR COBI OR RV2066 OR MT2126 OR MTCV49.05 OR MB2092.
OS Mycobacterium tuberculosis, and
OS Mycobacterium bovis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773, 1765;
RN [1]
RN SEQUENCE FROM N.A.
RC SPECIES=M.tuberculosis; STRAIN=H37RV;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Broach R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Fellwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrall B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
```

```

RL Nature 393:537-544 (1998).
RN [2]
RN SEQUENCE FROM N.A.
RC SPECIES=M.tuberculosis; STRAIN=CDC 1551 / Oshkosh;
RX MEDLINE=22206494; PubMed=12218036;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,
RA Kelowny J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;
RT "Whole-genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."
RL J. Bacteriol. 184:5479-5490 (2002).
RN [3]
RN SEQUENCE FROM N.A.
RC SPECIES=M.bovis; STRAIN=AF2122/97;
RX MEDLINE=22709107; PubMed=12788972;
RA Garnier T., Eiglmeier K., Camus J.-C., Medina N., Mansoor H.,
RA Pryor M., Duthoy S., Grondin S., Lacroix C., Monsemp C., Simon S.,
RA Harris B., Atkin R., Doggett J., Mayes R., Keating L., Wheeler P.R.,
RA Parkhill J., Barrall B.G., Cole S.T., Gordon S.V., Hewinson R.G.;
RT "The complete genome sequence of Mycobacterium bovis."
RL Proc. Natl. Acad. Sci. U.S.A. 100:7877-7882 (2003).
CC -!- FUNCTION: METHYLATES PRECORRIN-2 AT THE C-20 POSITION TO PRODUCE
CC PRECORRIN-3A (BY SIMILARITY).
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + precorrin-2 = S-
CC adenosyl-L-homocysteine + precorrin-3A.
CC -!- PATHWAY: Cobalamin biosynthesis.
CC -!- SIMILARITY: BELONGS TO A FAMILY THAT GROUPS SUMT, CYSG, CBIF/COBM
CC AND CBIL/COBI.
CC -----
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CC -----
DR EMBL; Z73966; CAA98214.1; -.
DR EMBL; AE007083; AAK46406.1; -.
DR EMBL; BX248341; CAD96945.1; -.
DR PIR; E70764; E70764.
DR TIGR; MT2126; -.
DR TuberculList; RV2066; -.
DR InterPro; IPR006364; Cobi_Cbil.
DR InterPro; IPR008363; CobiJ.
DR InterPro; IPR008078; Cor/por_Metransf.
DR InterPro; IPR003043; Uropor_Metransf.
DR Pfam; PF00590; TP_methylase; 2.
DR TIGRFAMs; TIGR01467; cobi_cbil; 1.
DR TIGRFAMs; TIGR01466; cobi_cbil; 1.
DR PROSITE; PS00839; SUMT_1; 1.
DR PROSITE; PS00840; SUMT_2; 1.
DR Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
DR Methyltransferase; Multifunctional enzyme; Complete proteome.
FT DOMAIN 1 243 PRECORRIN-2 C20-METHYLTRANSFERASE.
FT DOMAIN 244 508 PRECORRIN-3 METHYLASE.
SQ SEQUENCE 508 AA; 53910 MW; 95AC066F022C4DC1 CRC64;

Query Match 43.4%; Score 46; DB 1; Length 508;
Best Local Similarity 47.1%; Pred. No. 19;
Matches 8; Conservative 4; Mismatches 5; Indels 0; Gaps 0;

QY 1 MAGLPVVVGLSPGQEQY 17
DB 245 LTGVVVVGLGPGSDW 261

RESULT 6
CNA2_MOUSE
ID CNA2_MOUSE STANDARD; PRT; 1172 AA.
AC Q80XC6; Q8R3D7; Q99LT9;
```

DT 10-OCT-2003 (Rel. 42, Created)  
 DT 10-OCT-2003 (Rel. 42, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Protein C14orf102 homolog.  
 GN C14orf102.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 NCBI\_TaxID=10090;  
 [1]  
 RN CONCEPTUAL TRANSLATION, AND RECONSTRUCTION FROM EST.  
 RP Reynaud S.;  
 RA Unpublished observations (AUG-2003).  
 [2]  
 RP SEQUENCE OF 21-1172 FROM N.A.  
 RC TISSUE=Breast tumor, and Olfactory epithelium;  
 RX MEDLINE=22388257; PubMed=12477932;  
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,  
 RA Klausner R.D., Collins F.S., Wagner L., Sherman C.M., Schuler G.D.,  
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,  
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,  
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,  
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,  
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,  
 RA Raha S.S., Lequellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,  
 RA Bobak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,  
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,  
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,  
 RA Fahey J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,  
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,  
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,  
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,  
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalls D.E.,  
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;  
 RT "Generation and initial analysis of more than 15,000 full-length human  
 and mouse cDNA sequences.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).  
 CC -!- SIMILARITY: Contains 5 HAT repeats.  
 CC -!- CAUTION: This is a conceptual translation; a frameshift was  
 introduced in position 28 to produce the correct N-terminus, and  
 to extend the similarity with the human ortholog.  
 CC -!- CAUTION: Ref.1 (AAH51175) sequence differs from that shown due to  
 a stop codon in position 1146 which was translated as Gln to  
 extend the sequence.  
 CC -----  
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 CC -----  
 DR EMBL; BY182441; -; NOT ANNOTATED\_CDS.  
 DR EMBL; BC002230; AAH02230.1; -.  
 DR EMBL; BC025577; AAH25577.1; ALT\_INIT.  
 DR EMBL; BC051175; AAH51175.1; ALT\_SEQ.  
 DR InterPro; IPR003107; HAT.  
 DR SMART; SM00386; HAT; 5.  
 KW Repeat; Coiled coil.  
 FT DOMAIN 78 330 COILED COIL (POTENTIAL).  
 FT REPEAT 314 346 HAT 1.  
 FT REPEAT 404 436 HAT 2.  
 FT REPEAT 766 800 HAT 3.  
 FT REPEAT 986 1018 HAT 4.  
 FT REPEAT 1075 1109 HAT 5.  
 FT DOMAIN 86 93 POLY-LYS.  
 FT DOMAIN 146 154 POLY-ALA.  
 FT DOMAIN 707 707 R -> H (IN REF. 1; AAH25577).  
 FT CONFLICT 741 742 LS -> VC (IN REF. 1; AAH25577).  
 SQ SEQUENCE 1172 AA; 133424 MW; ABE37A8BCE7A528F CRC64;  
 Query Match 41.5%; Score 44; DB 1; Length 1172;  
 Best Local Similarity 53.3%; Pred. No. 94;  
 Matches 8; Conservative 1; Mismatches 6; Indels 0; Gaps 0;  
 QY 5 PAVVGLSPGQEVYHR 19  
 Db 351 PGYALGEGGEQEKHR 365  
 RESULT 7  
 MINT HUMAN  
 ID MINT\_HUMAN STANDARD; PRT: 3664 AA.  
 AC Q96T58; Q9H9A8; Q9NWH5; Q9UQ01; Q9Y556;  
 DT 10-OCT-2003 (Rel. 42, Created)  
 DT 10-OCT-2003 (Rel. 42, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Mx2-interacting protein (SMART/HDAC1 associated repressor protein).  
 GN MINT OR SHARP OR KIAA0929.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 NCBI\_TaxID=9606;  
 [1]  
 RP SEQUENCE FROM N.A., FUNCTION, SUBCELLULAR LOCATION, INDUCTION,  
 RP RNA-BINDING, AND INTERACTION WITH NCOR2; HDAC1; HDAC2; RBBP4; MED3;  
 RP RAR AND MTAIL1.  
 RP TISSUE=Liver, and Pituitary;  
 RC MEDLINE=21231190; PubMed=11331609;  
 RX Shi Y., Downes M., Xie W., Kao H.-Y., Ordentlich P., Tsai C.-C.,  
 RA Hon M., Evans R.M.;  
 RA "Sharp, an inducible cofactor that integrates nuclear receptor  
 repression and activation.";  
 RL Genes Dev. 15:1140-1151 (2001).  
 [2]  
 RP SEQUENCE FROM N.A.  
 RA Bird C.;  
 RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.  
 [3]  
 RP SEQUENCE OF 294-3664 FROM N.A.  
 RA Rhodes S., Huckle E.;  
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.  
 [4]  
 RP SEQUENCE OF 793-1595 FROM N.A., AND VARIANT PRO-1091.  
 RC TISSUE=Embryo, and Teratocarcinoma;  
 RA Isegai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,  
 RA Nishikawa T., Nagai K., Sugano S., Takahashi-Fujii A., Hara H.,  
 RA Tanase T., Nomura Y., Togiya S., Komai F., Hara R., Takeuchi K.,  
 RA Arita M., Nabekura T., Ishii S., Kawai Y., Saito K., Yamamoto J.,  
 RA Wakamatsu A., Nakamura Y., Nagahara K., Masuho Y., Oshima A.;  
 RL "NEDO human cDNA sequencing project.";  
 RT Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.  
 [5]  
 RP SEQUENCE OF 2002-3664 FROM N.A.  
 RC TISSUE=Brain;  
 RX MEDLINE=99246063; PubMed=10231032;  
 RA Nagase T., Ishikawa K.-I., Suyama M., Kikuno R., Hirose M.,  
 RA Miyajima N., Tanaka A., Kotani H., Nomura N., Ohara O.;  
 RT "Prediction of the coding sequences of unidentified human genes. XIII.  
 The complete sequences of 100 new cDNA clones from brain which code  
 for large proteins in vitro.";  
 RL DNA Res. 6:63-70 (1999).  
 [6]  
 RP INTERACTION WITH PPARD.  
 RX MEDLINE=21874127; PubMed=11867749;  
 RA Shi Y., Hoi M., Evans R.M.;  
 RT "The peroxisome proliferator-activated receptor delta, an integrator  
 of transcriptional repression and nuclear receptor signaling.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 99:2613-2618 (2002).  
 [7]  
 RP FUNCTION, AND INTERACTION WITH REPSUH.  
 RX MEDLINE=22261914; PubMed=12374742;  
 RA Oswald F., Kostezka U., Astrahantseff K., Bourteele S., Dillinger K.,  
 RA Zechner U., Ludwig L., Wilda M., Hameister H., Knoechel W., Lipray S.,  
 RA Schmid R.M.;



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DR EMBL; U67593; AAB99541.1; -.  
 DR PIR; A64490; A64490.  
 DR TIGR; MJ1522; -.

DR InterPro; IPR000878; Cor/por Metransf.  
 DR Pfam; PF005090; TP methylase; 1.  
 KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;  
 KW Methyltransferase; Complete proteome.  
 SQ SEQUENCE 211 AA; 23805 MW; 279A1A2B14369510 CRC64;

Query Match 40.6%; Score 43; DB 1; Length 211;  
 Best Local Similarity 54.5%; Pred. No. 23;  
 Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 7 VVGLSPGGEQY 17  
 :||: ||:|  
 Db 4 IVGIGFGDEY 14

## RESULT 9

MER METTI STANDARD; PRT; 326 AA.  
 AC Q9UKP0;  
 DT 15-MAR-2004 (Rel. 43, Created)  
 DT 15-MAR-2004 (Rel. 43, Last sequence update)  
 DT 15-MAR-2004 (Rel. 43, Last annotation update)  
 DE Coenzyme F420-dependent N(5),N(10)-methyltetrahydromethanopterin  
 DE reductase (EC 1.5.99.11) (Methylene-H(4)MPT reductase).  
 GN MER OR FFDA.  
 OS Methanobolus tindarius.  
 OC Archaea; Euryarchaeota; Methanomicrobia; Methanosarcinales;  
 OC Methanosarcinaceae; Methanobolus.  
 OX NCBI\_TaxID=2221;  
 RN [1]

## SEQUENCE FROM N.A.

RC STRAIN=DSM 2278;  
 RC MEDLINE=99132696; PubMed=9933933;  
 RA Wostenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,  
 RA Gottschalk G., Blaut M.;  
 RA "The F420H2-dehydrogenase from Methanobolus tindarius: cloning of the  
 RT ffd operon and expression of the genes in *Escherichia coli*.";  
 RL FEMS Microbiol. Lett. 170:389-398(1999).  
 CC -!- FUNCTION: Catalyzes the reversible reduction of methylene-H(4)MPT  
 CC to methyl-H(4)MPT (By similarity).  
 CC -!- CATALYTIC ACTIVITY: N(5),N(10)-methyltetrahydromethanopterin +  
 CC reduced coenzyme F420 = 5-methyl-5,6,7,8-tetrahydromethanopterin +  
 CC coenzyme F420.  
 CC -!- PATHWAY: Methanogenesis from carbon dioxide; fifth step.  
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).  
 CC -!- SIMILARITY: Belongs to the mer family.

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DR EMBL; AJ011519; CAB56639.1; -.  
 DR PIR; T45226; T45226.  
 DR HAMAP; MF\_01091; -.  
 DR InterPro; IPR002103; Bac\_luciferase.  
 DR Pfam; PF00296; Bac\_luciferase; 1.  
 KW Methanogenesis; One-carbon metabolism; Oxidoreductase.  
 SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;

Query Match 40.6%; Score 43; DB 1; Length 326;  
 Best Local Similarity 53.3%; Pred. No. 35;  
 Matches 8; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 MAGLPVVVGLSPGGEQ 15  
 :||: ||:|  
 Db 83 ISGGAUUGLPGGEQ 97

## RESULT 10

COBL MYCTU STANDARD; PRT; 390 AA.  
 AC Q1067L;  
 DT 15-JUL-1998 (Rel. 36, Created)  
 DT 15-DEC-1998 (Rel. 37, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Precorrin-6Y C5,15-methyltransferase [decarboxylating] (BC 2.1.1.132)  
 DE (Precorrin-6 methyltransferase) [Precorrin-6Y methylase].  
 GN COBL OR RV2072C OR MT2132 OR MTCV49.11C.  
 OS Mycobacterium tuberculosis.

OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
 OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.

OX NCBI\_TaxID=1773;  
 RN [1]

## SEQUENCE FROM N.A.

RC STRAIN=H37Rv;  
 RC MEDLINE=98295987; PubMed=9634230;  
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,  
 RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,  
 RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,  
 RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,  
 RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,  
 RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,  
 RA Ruster S., Seeger K., Skelton S., Squares S., Squares R.,  
 RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;  
 RT "Deciphering the biology of Mycobacterium tuberculosis from the  
 RL complete genome sequence.";  
 RL Nature 393:537-544(1998).  
 RN [2]

## SEQUENCE FROM N.A.

RC STRAIN=CDC 1551 / Oshkosh;  
 RC MEDLINE=22206494; PubMed=12218036;  
 RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,  
 RA Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,  
 RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,  
 RA Decher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,  
 RA Bishai W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;  
 RT "Whole-genome comparison of Mycobacterium tuberculosis clinical and  
 RT laboratory strains.";  
 RL J. Bacteriol. 184:5479-5490(2002).  
 CC -!- FUNCTION: CATALYZES THE METHYLATION OF BOTH C-5 AND C-15 IN  
 CC PRECORRIN-6Y TO FORM PRECORRIN-8X.  
 CC -!- CATALYTIC ACTIVITY: 2 S-adenosyl-L-methionine + precorrin-6Y = 2  
 CC S-adenosyl-L-homocysteine + precorrin-8X + CO(2).  
 CC -!- PATHWAY: Cobalamin biosynthesis.  
 CC -!- SIMILARITY: TO S-TYPHIMURIUM CBIE; ALSO, LOW, TO OTHER  
 CC METHYLASES INVOLVED IN COBALAMIN BIOSYNTHESIS.

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DR EMBL; Z73966; CAA98225.1; -.  
 DR EMBL; AE007063; AAK46412.1; -.  
 DR PIR; C70765; C70765.  
 DR TIGR; MT2132; -.  
 DR InterPro; IPR006365; Cobl.  
 DR InterPro; IPR000978; Cor/por Metransf.

```

DR InterPro; IPR000051; SAM bind.
DR Pfam; PF00590; TP_methylase; 1.
DR TIGRFAMs; TIGR01468; coBL_cblE; 1.
KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW Methyltransferase; Complete proteome.
FT CONFLICT 205 205 L -> P (IN REF. 2).
FT CONFLICT 327 327 L -> H (IN REF. 2).
SQ SEQUENCE 390 AA; 41854 MW; FB42EPF7562F21F3 CRC64;

Query Match 40.6%; Score 43; DB 1; Length 390;
Best Local Similarity 60.0%; Pred. No. 43;
Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 4 LPVAVGLSPGQEVYH 18
DB 55 LPVAVGLSPDGADLH 69

RESULT 11
YTSJ_CABEL STANDARD; PRT; 505 AA.
AC P90838;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Hypothetical UPF0027 protein F16A11.2 in chromosome I.
F16A11.2.
GN Caenorhabditis elegans.
OS Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
[1]
RN SEQUENCE FROM N.A.
RA Baynes C.;
RA Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
[2]
RN Durbini R.;
RA Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: Belongs to the UPF0027 (rtCB) family.
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-----
CC EMBL; Z81505; CAB04121.2; --
DR WormPep; F16A11.2; CE23663.
DR InterPro; IPR001233; UPF0027.
DR Pfam; PF01139; UPF0027; 1.
DR PROSITE; PS01288; UPF0027; 1.
KW Hypothetical protein.
SQ SEQUENCE 505 AA; 55230 MW; D528F702E2596909 CRC64;

Query Match 40.6%; Score 43; DB 1; Length 505;
Best Local Similarity 45.0%; Pred. No. 56;
Matches 9; Conservative 3; Mismatches 8; Indels 0; Gaps 0;

QY 1 MAGLPVAVGLSPGQEVYHRG 20
DB 72 VASLPGVGHSLGLPDIHSG 91

RESULT 12
C4AD_DROME STANDARD; PRT; 516 AA.
AC Q9V4T3;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)

```

10-OCT-2003 (Rel. 42, Last annotation update)  
 Probable cytochrome P450 4ad1 (EC 1.14.-.-) (CYPIVAD1).  
 CYP4AD1 OR CG2110.  
 Drosophila melanogaster (Fruit fly).  
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
 Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
 OC Ephydroidea; Drosophilidae; Drosophila.  
 NCBI\_TaxID=7227;  
 [1]  
 SEQUENCE FROM N.A.  
 STRAIN=Berkely;  
 MEDLINE=20196006; PubMed=10731132;  
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,  
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,  
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,  
 RA Sutton G.G., Wortman J.R., Vandeil M.D., Zhang Q., Chen L.X.,  
 RA Brandon R.C., Rogers Y.-H.C., Blazek R.G., Champe M., Pfeiffer B.D.,  
 RA Wan K.H., Doyle C., Baxter E.G., Helt J., Nelson C.R., Miklos G.L.G.,  
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfankoch C., Baldwin D.,  
 RA Ballwe R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,  
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,  
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,  
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,  
 RA Chertis J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,  
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,  
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,  
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,  
 RA Fessler C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,  
 RA Glodek A., Gong P., Gorrell J.H., Gu Z., Guan P., Harris M.,  
 RA Harris N.L., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,  
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,  
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,  
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,  
 RA Lasko P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.,  
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,  
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,  
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,  
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,  
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,  
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,  
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,  
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,  
 RA Swierkas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,  
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissenbach J.,  
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,  
 RA Ye J., Yeh R.-P., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,  
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Smith H.O.,  
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;  
 "The genome sequence of *Drosophila melanogaster*."  
 Science 287:2185-2195 (2000).  
 [2]  
 SEQUENCE FROM N.A.  
 STRAIN=Berkely; TISSUE=Embryo;  
 MEDLINE=22426066; PubMed=12537569;  
 RA Stapleton M., Carlson J.W., Brokstein P., Yu C., Champe M.,  
 RA George R.A., Guarini H., Kronmiller B., Pacleb J.M., Park S., Wan X.H.,  
 RA Rubin G.M., Celniker S.E.;  
 "A *Drosophila* full-length cDNA resource."  
 Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8 (2002).  
 RL -1- FUNCTION: May be involved in the metabolism of insect hormones and  
 CC in the breakdown of synthetic insecticides (By similarity).  
 CC -1- CATALYTIC ACTIVITY: RH + reduced flavoprotein + O(2) = ROH +  
 CC oxidized flavoprotein + H(2)O.  
 CC -1- SUBCELLULAR LOCATION: Membrane-bound. Endoplasmic reticulum  
 CC (Potential).  
 CC -1- SIMILARITY: Belongs to the cytochrome P450 family.  
 -----
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CC
CC EMBL; AE003837; AAF59092.1; -.
CC EMBL; AY061058; AAL28606.1; -.
CC HSSP; F14779; IJFZ.
CC FlyBase; FBgn0033292; Cyp4ad1.
CC InterPro; IPR001128; Cytochrome_P450.
CC Pfam; PF00067; P450; 1.
CC PRINTS; PR00385; P450.
CC PROSITE; PS00086; CYTOCHROME_P450; 1.
KW Oxidoreductase; Monooxygenase; Membrane; Heme; Microsome;
KW Endoplasmic reticulum; Hypothetical protein.
FT METAL 445 445 IRON (HEME AXIAL LIGAND) (BY SIMILARITY).
SQ SEQUENCE 516 AA; 58870 MW; 648EA22492AF58C7 CRC64;

Query Match 40.6%; Score 43; DB 1; Length 516;
Best Local Similarity 56.2%; Pred. No. 57;
Matches 9; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 4 LPVVVGLSPGQEQYHR 19
DB 469 LPVVVGLSPGQEQYHR 484

RESULT 13
CH60 PROAC STANDARD; PRT; 544 AA.
AC Q9K2U4;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE 60 kDa chaperonin (Protein Cpn60) (groEL protein) (Heat shock protein
DE 60).
GN GROEL OR GROEL OR HSP60.
OS Propionibacterium acnes.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Propionibacterineae; Propionibacteriaceae; Propionibacterium.
OX NCBI_TaxID=1747;
RN [1]
RP SEQUENCE FROM N.A.
RA Cho Y.;
RT "Propionibacterium acnes hsp60 gene.";
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=P37;
RA Farrar M.D., Ingham E., Holland K.T.;
RT "Cloning and sequencing of a groEL homolog from Propionibacterium
RT acnes.";
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Prevents misfolding and promotes the refolding and
CC proper assembly of unfolded polypeptides generated under stress
CC conditions (By similarity).
CC -!- SUBUNIT: Oligomer of 14 subunits composed of two stacked rings of
CC 7 subunits (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the chaperonin (HSP60) family.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC -----
CC EMBL; AB036414; BAA92382.1; -.
CC EMBL; AF222061; AAF33788.1; -.
CC HSSP; P06139; IGR.
CC HAMAP; MF_00600; -. 1.
CC InterPro; IPR001844; Chaperonin Cpn60.
CC InterPro; IPR002423; Cpn60/Tcf-1.
CC InterPro; IPR008950; GroEL-ATPase.

or send an email to license@isb-sib.ch).
DR PFam; PF00119; cpn60_TCP1; 1.
DR PRINTS; PR00298; CHAPERONIN60.
DR PROSITE; PR00304; TCOMPLEXTCP1.
DR PROSITE; PS00296; CHAPERONINS_CPN60; 1.
KW Chaperone; ATP-binding
SQ SEQUENCE 544 AA; 56839 MW; 9B4314A8BF8A94DC CRC64;

Query Match 40.6%; Score 43; DB 1; Length 544;
Best Local Similarity 52.9%; Pred. No. 60;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 1 MAGLPVVVGLSPGQEQY 17
DB 466 VAGLPAGQGLNAANDEY 482

RESULT 14
HNFA MOUSE STANDARD; PRT; 628 AA.
ID HNFA_MOUSE
AC P22361;
DT 01-AUG-1991 (Rel. 19, Created)
DT 01-AUG-1991 (Rel. 19, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Hepatocyte nuclear factor 1-alpha (HNF-1A) (Liver-specific
DE transcription factor 1F-B1) (LFB1).
GN TCF1 OR HNF1A OR HNF-1A OR HNF-1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=9108607; PubMed=2263635;
RA Kuo C.J., Conley P.B., Hsieh C.L., Francke U., Crabtree G.R.;
RT "Molecular cloning, functional expression, and chromosomal
RT localization of mouse hepatocyte nuclear factor 1.";
RL Proc. Natl. Acad. Sci. U.S.A. 87:9838-9842(1990).
CC -!- FUNCTION: Required for the expression of several liver specific
CC genes. Binds to the inverted palindrome 5'-GTTAATNATTAAC-3'.
CC -!- SUBUNIT: Binds DNA as a dimer.
CC -!- SUBCELLULAR LOCATION: Nuclear.
CC -!- TISSUE SPECIFICITY: Liver.
CC -!- SIMILARITY: Belongs to the HNF1 homeobox family.
CC -!- SIMILARITY: Contains 1 homeobox domain.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC
CC -----
CC EMBL; M57966; AAA37821.1; -.
CC PIR; A39262; A39262.
CC PDB; 1F93; 20-SEP-00.
CC PDB; 1G2Y; 17-JAN-01.
CC PDB; 1G2Z; 17-JAN-01.
CC PDB; 1G39; 17-JAN-01.
CC PDB; 1JB6; 11-JUL-01.
CC PDB; 1LFB; 31-OCT-93.
CC TRANSFAC; T01211; -.
CC MGD; MGI:98504; Tcf1.
CC GO; GO:0005634; C:nucleus; IDA.
CC InterPro; IPR006899; HNF-1_N.
CC InterPro; IPR006898; HNF1A_C.
CC InterPro; IPR001356; HNF1B_C.
CC InterPro; IPR006897; Homeobox.
CC Pfam; PF04814; HNF-1_N; 1.
CC Pfam; PF04813; HNF-1A_C; 1.
CC Pfam; PF04812; HNF-1B_C; 1.
CC SMART; SM00389; HOX; 1.
CC PROSITE; PS00027; HOMEBOX_1; 1.
```

DR PROSITE; PS50071; HOMEBOX 2; 1.  
 KW Transcription regulation; DNA-binding; Homeobox; Nuclear protein;  
 FT Activator; Trans-acting factor; 3D-structure.  
 FT DOMAIN 1 31 DIMERIZATION.  
 FT DOMAIN 71 80 ASP/GLU-RICH (ACIDIC) (POTENTIAL  
 FT INVOLVEMENT WITH TRANSCRIPTION).  
 FT NUCLEAR LOCALIZATION SIGNAL (POTENTIAL).  
 FT HOMEBOX.  
 FT DNA BIND 197 205  
 FT DNA BIND 199 279  
 FT DOMAIN 238 258 21 AMINO ACID LOOP BETWEEN HELIX 2 AND 3.  
 SQ SEQUENCE 628 AA; 67237 MW; 737920D1A369B9DD CRC64;

Query Match 40.6%; Score 43; DB 1; Length 628;  
 Best Local Similarity 57.1%; Pred. No. 70;  
 Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 MAGLEFAVVGSLSPGE 14  
 ||||| : : |||  
 404 MASLPQVMTIGPGE 417

Db

RESULT 15  
 HNFA\_RAT STANDARD; PRT; 628 AA.  
 ID HNFA\_RAT  
 AC P15257;  
 DT 01-APR-1990 (Rel. 14, Created)  
 DT 01-APR-1990 (Rel. 14, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Hepatocyte nuclear factor 1-alpha (HNF-1A) (liver-specific  
 DE Transcription factor LF-B1) (LFBI).  
 DE TCF1 OR HNF1A OR HNF-1A OR HNF-1.  
 OS Rattus norvegicus (Rat).  
 GN Rattus norvegicus (Rat).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
 OX NCBI\_TaxID=10116;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Liver;  
 RX MEDLINE=90003224; PubMed=2571419;  
 RA Frain M., Swart G., Monaci P., Nicosia A., Staempfli S., Frank R.,  
 RA Cortese R.;  
 RT "The liver-specific transcription factor LF-B1 contains a highly  
 RT diverged homeobox DNA binding domain.";  
 RL Cell 59:145-157(1989).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=91016926; PubMed=2216777;  
 RA Chouard T., Blumenfeld M., Bach I., Vandekerckhove J., Cereghini S.,  
 RA Yaniv M.;  
 RT "A distal dimerization domain is essential for DNA-binding by the  
 RT atypical HNF1 homeobox domain.";  
 RL Nucleic Acids Res. 18:5853-5863(1990).  
 RN [3]  
 RP SEQUENCE OF 166-628 FROM N.A.  
 RX MEDLINE=90249741; PubMed=1970973;  
 RA Baumhueter S., Mendel D.B., Conley P.B., Kuo C.J., Turk C.,  
 RA Graves M.K., Edwards C.A., Courtois G., Crabtree G.R.;  
 RT "HNF-1 shares three sequence motifs with the POU domain proteins and  
 RT is identical to LF-B1 and APF.";  
 RL Genes Dev. 4:372-379(1990).  
 RN [4]  
 RP SEQUENCE OF 1-12 FROM N.A.  
 RC STRAIN=Sprague-Dawley;  
 RA Tomei L., Piaggio G., Toniatti C., Lazzaro D., de Francesco R.,  
 RA Pozzi L., Gerstner J., Cortese R.;  
 RL Submitted (AUG-1992) to the EMBL/GenBank/DBJ databases.  
 RN [5]  
 RP POSITION OF HOMEBOX.  
 RX MEDLINE=90106643; PubMed=1967225;  
 RA Finney M.;  
 RT "The homeodomain of the transcription factor LF-B1 has a 21 amino  
 RT acid loop between helix 2 and helix 3.";  
 RL Cell 60:5-6(1990).  
 RN [6]

RP STRUCTURE BY NMR OF 1-32.  
 RX MEDLINE=91105074; PubMed=1988016;  
 RA Pastore A., de Francesco R., Barbato G., Castiglione Morelli M.A.,  
 RA Motta A., Cortese R.;  
 RT "1H resonance assignment and secondary structure determination of the  
 RT dimerization domain of transcription factor LFBI.";  
 RL Biochemistry 30:148-153(1991).  
 RN [7]  
 RP STRUCTURE BY NMR OF 195-286.  
 RX MEDLINE=93259120; PubMed=8491172;  
 RA Leitinger B., de Francesco R., Tomei L., Cortese R., Otting G.,  
 RA Wuethrich K.;  
 RT "The three-dimensional NMR-solution structure of the polypeptide  
 RT fragment 195-286 of the LFBI/HNF1 transcription factor from rat liver  
 RT comprises a nonclassical homeodomain.";  
 RL EMBO J. 12:1797-1803(1993).  
 RN [8]  
 RP STRUCTURE BY NMR OF 195-286.  
 RX MEDLINE=97272000; PubMed=9126845;  
 RA Schott O., Billeter M., Leitinger B., Wider G., Wuethrich K.;  
 RT "The NMR solution structure of the non-classical homeodomain from the  
 RT rat liver LFBI/HNF1 transcription factor.";  
 RL J. Mol. Biol. 267:673-683(1997).  
 RN [9]  
 RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS) OF 195-286.  
 RX MEDLINE=93259121; PubMed=8491173;  
 RA Ceska T.A., Tamers M., Monaci P., Nicosia A., Cortese R., Suck D.;  
 RT "The X-ray structure of an atypical homeodomain present in the rat  
 RT liver transcription factor LFBI/HNF1 and implications for DNA  
 RT binding.";  
 RL EMBO J. 12:1805-1810(1993).  
 CC -!- FUNCTION: Required for the expression of several liver specific  
 CC genes. Binds to the inverted palindrome 5'-GTTAATTATTAAAC-3'.  
 CC -!- SUBUNIT: Binds DNA as a dimer.  
 CC -!- SUBCELLULAR LOCATION: Nuclear.  
 CC -!- TISSUE SPECIFICITY: Liver.  
 CC -!- SIMILARITY: Belongs to the HNF1 homeobox family.  
 CC -!- SIMILARITY: Contains 1 homeobox domain.  
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 CC -----  
 CC EMBL; J03170; AAA41524.1; -;  
 CC EMBL; X54423; CAA38295.1; -;  
 CC EMBL; X67649; CAA47891.1; -;  
 CC EMBL; X53297; CAA37387.1; ALT\_INIT.  
 CC PIR; A33333; A33333.  
 CC PIR; S25485; S25485.  
 CC PDB; 1LFB; 31-OCT-93.  
 CC PDB; 2LFB; 12-MAR-97.  
 CC TRANSFAC; T00369; -;  
 CC InterPro; IPR006899; HNF-1\_N.  
 CC InterPro; IPR006898; HNF1A\_C.  
 CC InterPro; IPR006897; HNF1B\_C.  
 CC InterPro; IPR001356; Homeobox.  
 CC Pfam; PF04814; HNF-1\_N; 1.  
 CC Pfam; PF04813; HNF-1A\_C; 1.  
 CC Pfam; PF04812; HNF-1B\_C; 1.  
 CC SMART; SM00389; HOX; 1.  
 CC PROSITE; PS00027; HOMEBOX\_1; 1.  
 CC PROSITE; PS00071; HOMEBOX\_2; 1.  
 CC Transcription regulation; DNA-binding; Homeobox; Nuclear protein;  
 KW Activator; Trans-acting factor; 3D-structure.  
 FT DOMAIN 1 31 DIMERIZATION.  
 FT DOMAIN 71 80 ASP/GLU-RICH (ACIDIC) (POTENTIAL  
 FT INVOLVEMENT WITH TRANSCRIPTION).  
 FT DOMAIN 197 205 NUCLEAR LOCALIZATION SIGNAL (POTENTIAL).  
 FT DNA\_BIND 199 279 HOMEBOX.

FT DOMAIN 238 258 21 AMINO ACID LOOP BETWEEN HELIX 2 AND 3.  
 FT HELIX 208 218  
 FT TURN 219 220  
 FT HELIX 226 241  
 FT TURN 242 245  
 FT TURN 248 249  
 FT TURN 251 254  
 FT HELIX 255 257  
 FT HELIX 261 273  
 FT TURN 274 275  
 SQ SEQUENCE 628 AA; 67213 MW; 8D28099308C86A52 CRC64;  
 Query Match 40.6%; Score 43; DB 1; Length 628;  
 Best Local Similarity 57.1%; Pred. No. 70;  
 Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;  
 QY 1 MAGLPVVGLSPGE 14  
 Db ||||| : |||  
 404 MASLPQVMTIGFGE 417

Search completed: May 7, 2004, 12:34:31  
 Job time : 8.2 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:26:40 ; Search time 37.4 Seconds  
(without alignments)  
168.726 Million cell updates/sec

Title: US-09-786-214A-9

Perfect score: 106

Sequence: 1 MAGLPVVGLSPGEQYHRG 20

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: SP\_TREMBL 25;\*

2: SP\_bacteria;\*

3: SP\_fungi;\*

4: SP\_human;\*

5: SP\_invertebrate;\*

6: SP\_mammal;\*

7: SP\_mhc;\*

8: SP\_organelle;\*

9: SP\_phage;\*

10: SP\_plant;\*

11: SP\_rodent;\*

12: SP\_virus;\*

13: SP\_vertebrate;\*

14: SP\_unclassified;\*

15: SP\_rvirus;\*

16: SP\_bacteriap;\*

17: SP\_archaeap;\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	64	60.4	41	11	Q8k408 rattus norv
2	52	49.1	150	16	Q8PE55 xanthomonas
3	50	47.2	242	16	Q89Y25 bradyrhizob
4	50	47.2	277	16	Q9KQK6 vibrio chol
5	50	47.2	284	16	Q7VF15 helicobacte
6	50	47.2	381	16	Q9RL15 deinococcus
7	49	46.2	277	16	Q87RD3 vibrio para
8	49	46.2	4823	13	Q93321 fugu rubrip
9	48	45.3	192	15	Q8UT08 human immun
10	48	45.3	192	15	Q90D29 human immun
11	48	45.3	278	16	Q9CMF6 pasteurella
12	48	45.3	300	16	Q8EMZ7 oceanobacil
13	48	45.3	485	10	Q7XNE4 oryza sativ
14	47	44.3	192	15	Q994Q2 human immun
15	47	44.3	192	15	Q9QSR2 human immun
16	47	44.3	192	15	Q994B2 human immun

17	47	44.3	250	16	Q8D0G6 versinia pe
18	47	44.3	280	16	Q8ZP42 salmonella
19	47	44.3	280	16	Q8XDC4 escherichia
20	47	44.3	280	16	Q8Z7F3 salmonella
21	47	44.3	289	16	Q8KRG2 chlorobium
22	47	44.3	293	16	O25975 helicobacte
23	47	44.3	293	16	Q9ZJ12 helicobacte
24	47	44.3	390	16	Q98CK1 rhizobium l
25	47	44.3	661	10	Q8S8J8 arabidopsis
26	46.5	43.9	242	16	Q7WCX1 bordetella
27	46.5	43.9	298	16	Q7WSD6 bordetella
28	46.5	43.9	298	16	Q7VTI2 bordetella
29	46.5	43.9	563	16	Q9X8S9 streptomyce
30	46	43.4	140	16	Q8P5G3 xanthomonas
31	46	43.4	143	4	Q8NAW2 homo sapien
32	46	43.4	192	15	Q90CX5 human immun
33	46	43.4	192	15	Q8UTJ5 human immun
34	46	43.4	192	15	O70887 human immun
35	46	43.4	274	16	Q9PPC9 campylobact
36	46	43.4	300	16	O34990 bacillus su
37	46	43.4	462	16	Q9KZB0 streptomyce
38	46	43.4	753	16	Q89T31 bradyrhizob
39	45	42.5	85	6	O77786 canis fami
40	45	42.5	192	15	Q90ML0 human immun
41	45	42.5	192	15	Q994N4 human immun
42	45	42.5	192	15	Q91W51 human immun
43	45	42.5	192	15	Q9WQH9 human immun
44	45	42.5	192	15	Q9WFP9 human immun
45	45	42.5	192	15	Q99BN6 human immun

## ALIGNMENTS

### RESULT 1

Q8K408 PRELIMINARY; PRT; 41 AA.

AC Q8K408, 2002 (Tremblrel. 22, Created)  
DT 01-OCT-2002 (Tremblrel. 22, last sequence update)  
DT 01-OCT-2002 (Tremblrel. 22, last annotation update)  
DE Truncated macrophage colony stimulating factor.  
GN CSF1.

OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
OX NCBI\_TaxID=10116;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=LEW.tl;  
RX MEDLINE=22069908; PubMed=12074592;  
RA Dobbins D.E., Sood R., Hashimoto A., Hansen C.T., Wilder R.L.,  
RA Remmers E.F.;  
RT "Mutation of macrophage colony stimulating factor (CSF1) causes  
osteopetrosis in the tl rat."

RL Biochem. Biophys. Res. Commun. 294:1114-1120(2002).

DR EMBL; AF514357; AAME4137.1; -

SQ SEQUENCE 41 AA; 4178 MW; 1D342C19BD18AA41 CRC64;

Query Match 60.4%; Score 64; DB 11; Length 41;  
Best Local Similarity 70.0%; Pred. No. 0.014;  
Matches 14; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 MAGLPVVGLSPGEQYHRG 20

DB 18 MDGLPAAAGLSPREQCCRG 37

### RESULT 2

Q8PE55 PRELIMINARY; PRT; 150 AA.

ID Q8PE55

AC Q8PE55; 01-OCT-2002 (Tremblrel. 22, Created)

DR Pfam; PF03737; Methyltransf\_6; 1.  
KW Complete proteome.  
SQ SEQUENCE 242 AA; 25216 MW; 588F9COB9396414B CRC64;

Query Match 47.2%; Score 50; DB 16; Length 242;  
Best Local Similarity 55.0%; Pred.No.14;  
Matches 11; Conservative 1; Mismatches 4; Indels 4; Gaps 1;

QY 1 MAGLPAVVGLSPGGEQYHRG 20  
: ||| ||||| |||||  
DB 63 LARLPGAVGLKP----YHRG 78  
: ||| ||||| |||||

RESULT 4  
Q9KQK6 PRELIMINARY; PRT; 277 AA.  
ID Q9KQK6  
AC Q9KQK6  
DT 01-OCT-2000 (T-EMBLrel. 15, Created)  
DT 01-OCT-2000 (T-EMBLrel. 15, Last sequence update)  
DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)  
DE Formyltetrahydrofolate deformylase.  
GN VC1992.  
OS Vibrio cholerae.  
OC Bacteria; Proteobacteria; Gammaproteobacteria; Vibrionales;  
OC Vibrionaceae; Vibrio.  
CX NCBI\_TaxID=666;  
[1]  
RN SEQUENCE FROM N.A.  
RP STRAIN=El Tor N16961 / Serotype O1;  
RC MEDLINE=20406833; PubMed=10952301;  
RA Heidelberg J.F., Elsen J.A., Nelson W.C., Clayton R.A., Gwinn M.L.,  
RA Dodson R.J., Haft D.H., Hickey E.K., Peterson J.D., Umayam I.A.,  
RA Gill S.R., Nelson K.E., Read T.D., Tettelin H., Richardson D.,  
RA McDonalad M.D., Vamathevan J., Bass S., Qin H., Dragoi I., Sellers P.,  
RA Emondale T., Uterback T., Fleischmann R.D., Nierman W.C., White O.,  
RA Salzberg S.L., Smith H.O., Colwell R.R., Mekalanos J.J., Venter J.C.,  
RA Fraser C.M.;  
RT "DNA sequence of both chromosomes of the cholera pathogen Vibrio  
cholerae.";  
RL Nature 406:477-483(2000).  
DR EMBL; AE004274; AAF95140.1; -.  
DR PIR; F82130; F82130.  
DR HGSP; P08179; 2GAR.  
DR TIGR; VC1992; -.  
DR GO; GO:0016597; F:amino acid binding; IEA.  
DR GO; GO:0008864; F:formyltetrahydrofolate deformylase activity; IEA.  
DR GO; GO:0016742; F:hydroxymethyl-, formyl- and related transfe. .; IEA.  
DR GO; GO:0006189; P:de novo IMP biosynthesis; IEA.  
DR GO; GO:0009058; P:biosynthesis; IEA.  
DR InterPro; IPR002912; ACT.  
DR InterPro; IPR002376; formyl\_transf.  
DR InterPro; IPR004810; PurU.  
DR Pfam; PF01842; ACT; 1.  
DR Pfam; PF00551; formyl\_transf; 1.  
DR PRINTS; PR01575; PF44HYDLASE.  
DR TIGRFAMs; TIGR00655; PurU; 1.  
KW Complete proteome.  
SQ SEQUENCE 277 AA; 31373 MW; A703491654753DC6 CRC64;

Query Match 47.2%; Score 50; DB 16; Length 277;  
Best Local Similarity 52.9%; Pred.No.16;  
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0

QY 4 LPVVVGLSPGGEQYHRG 20  
||| :||| :|||  
DB 190 LPAFIGAKPYQQAVERG 206  
||| :||| :|||

RESULT 5  
Q7VFI5 PRELIMINARY; PRT; 284 AA.  
ID Q7VFI5  
AC Q7VFI5  
DT 01-OCT-2003 (T-EMBLrel. 25, Created)

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DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Formyltetrahydrofolate deformylase PurU (BC 3.5.1.10).
GN PUKU OR HH1691.
OS Helicobacter hepaticus.
OC Bacteria; Proteobacteria; Epsilonproteobacteria; Campylobacteriales;
OC Helicobacteraceae; Helicobacter.
OX NCBI_TaxID=32025;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 51449 / 3B1;
RX MEDLINE=22709201; PubMed=12810954;
RA Suerbaum S., Josenhans C., Stenzenbach T., Drescher B., Brandt P.,
RA Bell M., Droege M., Fartmann B., Fischer H.-P., Ge Z., Hoerster A.,
RA Holland R., Klein K., Koenig J., Macko L., Mendz G.L., Nyakatura G.,
RA Schauer D.B., Shen Z., Weber J., Frosch M., Fox J.G.;
RT "The complete genome sequence of the carcinogenic bacterium
RT Helicobacter hepaticus."
RL Proc. Natl. Acad. Sci. U.S.A. 100:7901-7906(2003).
RW EMBL; AE017149; AAP78288.1; -.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 284 AA; 32660 MW; AAED2DC5086236E3 CRC64;

Query Match 47.2%; Score 50; DB 16; Length 284;
Best Local Similarity 52.9%; Pred. No. 17;
Matches 9; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 4 LPVVGLSPGEQYHRG 20
DB 197 LPAFIGANPYQAYERG 213

RESULT 6
Q9RRL5 ID Q9RRL5 PRELIMINARY; PRT; 381 AA.
AC Q9RRL5
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Cytochrome P450.
DN DR2473.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
RX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Ueberback T., Zalewski C.,
RA Makarova K.S., Aravind L., Daly M.J., Manton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RT "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1."
RL Science 286:1571-1577(1999).
CC -!- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.
DR EMBL; AE002076; AAF12016.1; -.
DR PIR; F75270; F75270.
DR TIGR; DR2473; -.
DR GO; GO:0004497; F:monoxygenase activity; IEA.
DR DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR001128; Cytochrome_P450.
DR Pfam; PF00067; p450; 1.
DR PRINTS; PR00385; P450.
DR PROSITE; PS00086; CYTOCHROME P450; 1.
KW Heme; Monoxygenase; Oxidoreductase; Complete proteome.
SQ SEQUENCE 381 AA; 41940 MW; F191EA69F1797B53 CRC64;

Query Match 47.2%; Score 50; DB 16; Length 381;

Best Local Similarity 100.0%; Pred. No. 23;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GLPAVVGSLSP 12
DB 51 GLPAVVGSLSP 60

RESULT 7
Q87RD3 ID Q87RD3 PRELIMINARY; PRT; 277 AA.
AC Q87RD3
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Formyltetrahydrofolate deformylase.
DN VP0864.
OS Vibrio parahaemolyticus.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Vibrionales;
OC Vibrionaceae; Vibrio.
OX NCBI_TaxID=670;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R1MD 2210633 / Serotype O3:K6;
RX MEDLINE=22508454; PubMed=12620739;
RA Makino K., Oshima K., Kurokawa K., Yokoyama K., Uda T., Tagomori K.,
RA Iijima Y., Najima M., Nakano M., Yamashita A., Kubota Y., Kimura S.,
RA Yasunaga T., Honda T., Shinagawa H., Hattori M., Iida T.;
RT "Genome sequence of Vibrio parahaemolyticus: a pathogenic mechanism
RT distinct from that of V. cholerae."
RL Lancet 361:743-749(2003).
RW EMBL; AP005075; BAC59127.1; -.
DR GO; GO:0016597; F:amino acid binding; IEA.
DR GO; GO:0008864; F:formyltetrahydrofolate deformylase activity; IEA.
DR GO; GO:0016742; F:hydroxymethyl-, formyl- and related transfe. .; IEA.
DR GO; GO:0006189; P:de novo IMP biosynthesis; IEA.
DR GO; GO:0009058; P:biosynthesis; IEA.
DR InterPro; IPR002912; ACT.
DR InterPro; IPR002376; formyl_transf.
DR InterPro; IPR004810; PurU.
DR Pfam; PF01842; ACT; 1.
DR Pfam; PF00551; formyl_transf; 1.
DR PRINTS; PR01575; FFH4HYDLASE.
KW Complete proteome.
SQ SEQUENCE 277 AA; 31620 MW; D5F0712E1537CF4C CRC64;

Query Match 46.2%; Score 49; DB 16; Length 277;
Best Local Similarity 52.9%; Pred. No. 23;
Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPVVGLSPGEQYHRG 20
DB 190 LPAFIGAKPYQAYDRG 206

RESULT 8
Q93321 ID Q93321 PRELIMINARY; PRT; 4823 AA.
AC Q93321
DT 01-NOV-1998 (TrEMBLrel. 08, Created)
DT 01-NOV-1998 (TrEMBLrel. 08, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE All-1 related protein.
DN ALR.
OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_TaxID=31033;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=99177347; PubMed=10077531;

```

RA Gellner K., Brenner S.;  
 RT "Analysis of 148 kb of genomic DNA around the wnt1 locus of Fugu  
 rubripes."; 9:251-258(1999).  
 RL Genome Res. 9:251-258(1999).  
 CC -1- SIMILARITY: CONTAINS 1 SET DOMAIN.  
 DR EMBL; AF056116; AAC34383.1; -.  
 DR GO; GO:0005634; C:nucleus; IEA.  
 DR GO; GO:0003677; F:DNA binding; IEA.  
 DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.  
 DR InterPro; IPR003889; FYrich\_C.  
 DR InterPro; IPR003888; FYrich\_N.  
 DR InterPro; IPR000910; HMG 12 box.  
 DR InterPro; IPR003616; PostSET.  
 DR InterPro; IPR001214; SET.  
 DR InterPro; IPR001965; Znf PHD.  
 DR InterPro; IPR001841; Znf\_ring.  
 DR Pfam; PF00628; PHD; 5.  
 DR Pfam; PF00856; SET; 1.  
 DR SMART; SM00542; FYRC; 1.  
 DR SMART; SM00541; FYRN; 1.  
 DR SMART; SM00398; HMG; 1.  
 DR SMART; SM00249; PHD; 8.  
 DR SMART; SM00508; PostSET; 1.  
 DR SMART; SM00184; RING; 4.  
 DR SMART; SM00317; SET; 1.  
 DR PROSITE; PS0868; POST SET; 1.  
 DR PROSITE; PS0280; SET; 1.  
 DR PROSITE; PS01359; ZF PHD 1; 1.  
 DR PROSITE; PS00016; ZF PHD 2; 1.  
 DR PROSITE; PS00089; ZF RING 2; 1.  
 DR PROSITE; PS00089; ZF RING 2; 1.  
 SQ SEQUENCE 4823 AA; 526260 MW; BD0C5F4EAD0F9C07 CRC64;

Query Match 46.2%; Score 49; DB 13; Length 4823;  
 Best Local Similarity 66.7%; Pred. No. 5.2e+02;  
 Matches 10; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 5 PAVVGLSPGQEQYHR 19  
 DB 1900 PALGGLSPELEKHR 1914  
 ||| ||| ||| |||

RESULT 9  
 Q8UTQ8 PRELIMINARY; PRT; 192 AA.  
 ID Q8UTQ8  
 AC Q8UTQ8  
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)  
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Vif protein (Virion infectivity factor) (SOR protein).  
 GN VIF.  
 OS Human immunodeficiency virus 1.  
 OC Viruses; Retrovird viruses; Retroviridae; Lentivirus.  
 OX NCBI\_TaxID=11676;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=98BWC14.a3;  
 RA Novitsky V.A., Smith U.R., Gilbert P., McLane M.F., Chigwedere P.,  
 RA Williamson C., Ndung'u T., Klein I., Chang S.-Y., Peter T., Thior I.,  
 RA Foley B.T., Gaolekwe S., Rybak N., Gaseitsiwe S., Vannberg F.,  
 RA Marlink R., Lee T.-H., Essex M.  
 RA "HIV-1 subtype C molecular phylogeny: consensus sequence for an AIDS  
 RT vaccine design."  
 RT Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.  
 RL -1- FUNCTION: DETERMINES VIRUS INFECTIVITY (BY SIMILARITY).  
 CC EMBL; AF443078; AAL34591.1; -.  
 DR GO; GO:0019058; P:viral infectious cycle; IEA.  
 DR InterPro; IPR000475; Viral\_infect.  
 DR Pfam; PF00559; Vif; 1.  
 DR PRINTS; PR00349; VIRIONINFECT.  
 DR ProDom; PD000063; Viral\_infect; 1.  
 DR HSSP; P08179; 1GRG.  
 DR GO; GO:0016597; F:amino acid binding; IEA.

Query Match 45.3%; Score 48; DB 15; Length 192;  
 Best Local Similarity 66.7%; Pred. No. 22;  
 Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 9 GLSPGQEQYHRG 20  
 DB 71 GLQPGEREWHLG 82  
 ||| ||| ||| |||

RESULT 10  
 Q90D29 PRELIMINARY; PRT; 192 AA.  
 ID Q90D29  
 AC Q90D29  
 DT 01-DEC-2001 (TrEMBLrel. 19, Created)  
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Vif protein (Virion infectivity factor) (SOR protein).  
 GN VIF.  
 OS Human immunodeficiency virus 1.  
 OC Viruses; Retrovird viruses; Retroviridae; Lentivirus.  
 OX NCBI\_TaxID=11676;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=97TZ01;  
 RX MEDLINE=21395692; PubMed=11504977;  
 RA Hoelscher M., Kim B., Maboko L., Mhalu F., von Sonnenburg F.,  
 RA Birx D.L., McCutchan F.E.,  
 RA the UNAIDS Network for HIV Isolation Characterization.;  
 RT "High proportion of unrelated HIV-1 intersubtype recombinants in the  
 RT Mbeya region of southwest Tanzania.";  
 RL AIDS 15:1461-1470(2001).  
 CC -1- FUNCTION: DETERMINES VIRUS INFECTIVITY (BY SIMILARITY).  
 DR EMBL; AF361871; AAK94212.1; -.  
 DR GO; GO:0019058; P:viral infectious cycle; IEA.  
 DR InterPro; IPR000475; Viral\_infect.  
 DR Pfam; PF00559; Vif; 1.  
 DR PRINTS; PR00349; VIRIONINFECT.  
 DR ProDom; PD000063; Viral\_infect; 1.  
 DR AIDS.  
 SQ SEQUENCE 192 AA; 22674 MW; 11B799C5DEA99F77 CRC64;

Query Match 45.3%; Score 48; DB 15; Length 192;  
 Best Local Similarity 66.7%; Pred. No. 22;  
 Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 9 GLSPGQEQYHRG 20  
 DB 71 GLQPGEREWHLG 82  
 ||| ||| ||| |||

RESULT 11  
 Q9CMF6 PRELIMINARY; PRT; 278 AA.  
 ID Q9CMF6  
 AC Q9CMF6  
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)  
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE PurU.  
 GN PURU OR PM0873.  
 OS Pasteurella multocida.  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;  
 OC Pasteurellaceae; Pasteurella.  
 OX NCBI\_TaxID=747;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=Pm70;  
 RA May B.J., Zhang Q., Li L.I., Paustian M.L., Whittam T.S., Kapur V.;  
 RA "Complete genomic sequence of Pasteurella multocida Pm70.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 98:3460-3465(2001).  
 DR EMBL; AE006126; AAK02957.1; -.  
 DR HSSP; P08179; 1GRG.  
 DR GO; GO:0016597; F:amino acid binding; IEA.

DR GO: GO:000864; F:formyltetrahydrofolate deformylase activity; IEA.  
 DR GO: GO:0016742; F:hydroxymethyl-, formyl- and related transfe. . .; IEA.  
 DR GO: GO:0006189; P:'de novo' IMP biosynthesis; IEA.  
 DR GO: GO:0009058; P:biosynthesis; IEA.  
 DR InterPro: IPR002912; ACT.  
 DR InterPro: IPR002376; formyl\_transf.  
 DR InterPro: IPR004810; PurU.  
 DR Pfam: PF01842; ACT; 1.  
 DR Pfam: PF00551; formyl transf; 1.  
 DR PRINTS: PR01575; FFH4HYDLASE.  
 DR TIGRfams: TIGR00655; PurU; 1.  
 DR Complete proteome.  
 SK SEQUENCE 278 AA; 32086 MW; F303E499DFFA0B70 CRC64;  
 Query Match 45.3%; Score 48; DB 16; Length 278;  
 Best Local Similarity 52.9%; Pred. No. 33;  
 Matches 9; Conservative 1; Mismatches 7; Indels 0; Gaps 0;  
 QY 4 LPVVGLSPGQEQYHRG 20  
 Db 191 LPAFIGAKPYHQAYERG 207  
 RESULT 12  
 Q8EMZ7 PRELIMINARY; PRT; 300 AA.  
 AC Q8EMZ7  
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)  
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Formyltetrahydrofolate deformylase (EC 3.5.1.10).  
 GN OB2693.  
 OS Oceanobacillus iheyensis.  
 OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Oceanobacillus.  
 OX NCBI\_TaxID=182710;  
 RN [1]  
 SEQUENCE FROM N.A.  
 RC STRAIN=HTE831 / DSM 14371 / JCM 11309;  
 RX MEDLINE=22220767; PubMed=12235376;  
 RA Takami H., Takaki Y., Uchiyama I.;  
 RT "Genome sequence of Oceanobacillus iheyensis isolated from the Iheya  
 RT Ridge and its unexpected adaptive capabilities to extreme  
 RT environments.";  
 RL Nucleic Acids Res. 30:3927-3935(2002).  
 DR EMBL; AF004602; BAC14649.1; -  
 DR GO: GO:0016597; F:amino acid binding; IEA.  
 DR GO: GO:000864; F:formyltetrahydrofolate deformylase activity; IEA.  
 DR GO: GO:0016787; F:hydrolase activity; IEA.  
 DR GO: GO:0016742; F:hydroxymethyl-, formyl- and related transfe. . .; IEA.  
 DR GO: GO:0006189; P:'de novo' IMP biosynthesis; IEA.  
 DR GO: GO:0009058; P:biosynthesis; IEA.  
 DR InterPro: IPR002912; ACT.  
 DR InterPro: IPR002376; formyl\_transf.  
 DR InterPro: IPR004810; PurU.  
 DR Pfam: PF01842; ACT; 1.  
 DR Pfam: PF00551; formyl transf; 1.  
 DR PRINTS: PR01575; FFH4HYDLASE.  
 DR TIGRfams: TIGR00655; PurU; 1.  
 DR Hydrolase; Complete proteome.  
 KW SEQUENCE 300 AA; 34894 MW; D44D0F0CC596BF2 CRC64;  
 Query Match 45.3%; Score 48; DB 16; Length 300;  
 Best Local Similarity 52.9%; Pred. No. 36;  
 Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;  
 QY 4 LPVVGLSPGQEQYHRG 20  
 Db 212 LPAFIGAKPYERAYDRG 228  
 RESULT 13  
 Q7XNE4 PRELIMINARY; PRT; 485 AA.  
 ID Q7XNE4

AC Q7XNE4;  
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE OSUNBA0009P12.18 protein.  
 GN OSUNBA0009P12.18.  
 OS Oryza sativa (Rice).  
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
 OC Ehrhartoideae; Oryzae; Oryza.  
 OX NCBI\_TaxID=4530;  
 RN [1]  
 SEQUENCE FROM N.A.  
 RA Liu B., Feng Q., Huang Y.C., Li Y., Zhu J.J., Zhao Q., Hu X.,  
 RA Liu Y.L., Mu J., Yu Z., Chen L., Fan D.L., Weng Q.J., Zhang L.,  
 RA Lu Y.Q., Yu S.L., Liu X.H., Lu T.T., Zhang Y.J., Lu Y., Li C., Li T.,  
 RA Zhang Y., Hu H., Jia P.X., Qian Y.M., Yang K., Zhou B., Chen Z.H.,  
 RA Hao P., Zhang L., Wu M., Zhang R.Q., Guan J.P., Fu G., Wang S.Y.,  
 RA Ren S.X., Lv G., Lin W., Gu W.Q., Zhu G.F., Tu Y.F., Jia J., Yin H.F.,  
 RA Zhang Y., Cai Z., Chen J., Kang H., Chen X.Y., Shao C.Y., Sun Y.,  
 RA Hu Q.P., Zhang X.L., Zhang W., Wang L.J., Ding C.W., Sheng H.H.,  
 RA Gu J.L., Chen S.T., Ni L., Zhu F.H., Hong G.F.;  
 RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AL662952; CAE04133.1; -  
 SQ SEQUENCE 485 AA; 52334 MW; F7685952D5538456 CRC64;  
 Query Match 45.3%; Score 48; DB 10; Length 485;  
 Best Local Similarity 60.0%; Pred. No. 61;  
 Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;  
 QY 4 LPVVGLSPGQEQYH 18  
 Db 317 LPALSKLSPGQAAHYH 331  
 RESULT 14  
 Q994Q2 PRELIMINARY; PRT; 192 AA.  
 ID Q994Q2  
 AC Q994Q2;  
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)  
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Vif protein (Viron infectivity factor) (SOR protein).  
 GN VIF.  
 OS Human immunodeficiency virus 1.  
 OC Viruses; Retroid viruses; Retroviridae; Lentivirus.  
 OX NCBI\_TaxID=11676;  
 RN [1]  
 SEQUENCE FROM N.A.  
 RC STRAIN=96ZM753;  
 RX MEDLINE=21094715; PubMed=11177395;  
 RA Rodenburg C.M., Li Y., Trask S.A., Chen Y., Decker J., Robertson D.L.,  
 RA Kalish M.L., Shaw G.M., Allen S., Hahn B.H., Gao F.;  
 RT "Near full-length clones and reference sequences for subtype C  
 RT isolates for HIV type 1 from three different continents.";  
 RL AIDS Res. Hum. Retroviruses 17:161-168(2001).  
 RN [2]  
 SEQUENCE FROM N.A.  
 RC STRAIN=96ZM751;  
 RA Rodenburg C.M., Li Y., Trask S.A., Chen Y., Decker J., Robertson D.L.,  
 RA Allen S., Shaw G.M., Hahn B.H., Gao F.;  
 RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.  
 CC -I- FUNCTION: DETERMINES VIRUS INFECTIVITY (BY SIMILARITY).  
 DR EMBL; AF286225; AAK30974.1; -  
 DR GO: GO:0019058; P:viral infectious cycle; IEA.  
 DR InterPro: IPR000475; Viral\_infect.  
 DR Pfam: PF00559; Vif; 1.  
 DR PRINTS: PR00349; VIRIONINFECT.  
 DR Prodrom; PD000063; Viral\_infect; 1.  
 KW AIDS.  
 SQ SEQUENCE 192 AA; 22727 MW; 55E01D4BBBCD93DC6 CRC64;  
 Query Match 44.3%; Score 47; DB 15; Length 192;



Best Local Similarity 66.7%; Pred. No. 32;  
Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 9 GLSPGGEYHRG 20  
|||:|:|  
Db 71 GLHFGEREHLG 82

## RESULT 15

Q9QSR2 PRELIMINARY; PRT; 192 AA.  
AC Q9QSR2;  
DT 01-MAY-2000 (TREMELrel. 13, Created)  
DT 01-MAY-2000 (TREMELrel. 13, Last sequence update)  
DT 01-JUN-2003 (TREMELrel. 24, Last annotation update)  
DE Vif protein (Vifon infectivity factor) (SOR protein).  
GN VIF.  
OS Human immunodeficiency virus 1.  
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=V1850;  
RX MEDLINE=20192166; PubMed=10725202;  
RA Laukkanen T., Carr J.K., Janssens W., Liitsola K., Gotte D.,  
RA McCutchan F.E., Op de Coul E., Cornelissen M., Heyndrickx L.,  
RA van der Groen G., Salminen M.O.;  
RT "Virtually full-length subtype F and F/D recombinant HIV-1 from Africa  
and South America.";  
RL Virology 269:95-104(2000).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC STRAIN=V1850;  
RA Carr J.K., Kim B., Sanders-Buell E., Salminen M.O., Alaeus A.,  
RA Albert J.A., Birx D.L., McCutchan F.E.;  
RT "HIV-1 isolate V1850 from Zaire, complete genome.";  
RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.  
CC -!- FUNCTION: DETERMINES VIRUS INFECTIVITY (BY SIMILARITY).  
DR EMBL; AF077336; AAD46089.1; -;  
DR GO; GO:0019058; P:Viral infectious cycle; IEA.  
DR InterPro; IPR000475; Viral\_infect.  
DR Pfam; PF00559; Vif; 1.  
DR PRINTS; PR00349; VIRIONINFECT.  
DR ProDom; PD000063; Viral\_infect; 1.  
KW AIDS.  
SQ SEQUENCE 192 AA; 22629 MW; F165805BFCDA4427 CRC64;

Query Match 44.3%; Score 47; DB 15; Length 192;  
Best Local Similarity 66.7%; Pred. No. 32;  
Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 9 GLSPGGEYHRG 20  
|||:|:|  
Db 71 GLHFGEREHLG 82

Search completed: May 7, 2004, 12:37:52  
Job time : 39.5667 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 55.8 Seconds  
(without alignments)  
101.272 Million cell updates/sec

Title: US-09-786-214A-9

Perfect score: 106

Sequence: 1 MAGLPAAVVLSPGQGEYHRG 20

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A Geneseq 29Jan04:\*

1: geneseqp1980s:\*

2: geneseqp1990s:\*

3: geneseqp2000s:\*

4: geneseqp2001s:\*

5: geneseqp2002s:\*

6: geneseqp2003as:\*

7: geneseqp2003bs:\*

8: geneseqp2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	106	100.0	20	3	AAy84265 Truncated
2	106	100.0	25	3	AAy84264 Peptide o
3	75	70.8	15	3	AAy84269 Peptide d
4	72	67.9	14	3	AAy84266 Peptide d
5	68	64.2	13	3	AAy84267 Peptide d
6	65	61.3	13	3	AAy84268 Peptide d
7	50	47.2	277	6	ABu49441 Protein e
8	49	46.2	191	3	AAy84265 HIV-1 non
9	49	46.2	278	6	ABu30626 Protein e
10	48	45.3	278	6	ABU39195 Protein e
11	48	45.3	287	6	ABu17217 Protein e
12	48	45.3	303	6	ADa36829 Acinetoba
13	47	44.3	151	5	ABu50765 Helicobac
14	47	44.3	155	5	ABu51750 Helicobac
15	47	44.3	192	3	AAy84269 HIV-1 non
16	47	44.3	238	6	ABu27483 Protein e
17	47	44.3	278	6	ABu44771 Protein e
18	47	44.3	280	6	ABu48054 Protein e
19	47	44.3	280	6	ABU31449 Protein e
20	47	44.3	280	6	ABU15054 Protein e
21	47	44.3	280	6	ABU47098 Protein e
22	47	44.3	282	6	ABU41187 Protein e
23	47	44.3	293	6	ABM69037 Photorhab
24	47	44.3	293	2	AAW98485 H. pylori
25	47	44.3	293	6	ABU31073 Protein e

26	46	43.4	143	5	AAO21677 Human sec
27	46	43.4	143	7	ADB64633 Human pro
28	46	43.4	187	6	ABu21296 Protein e
29	46	43.4	192	3	AAy84265 HIV-1 non
30	46	43.4	234	4	AAy84268 Human imm
31	46	43.4	274	6	ABu26498 Protein e
32	46	43.4	300	2	AAy16108 A formate
33	46	43.4	300	2	ABu24038 Novel hum
34	45	42.5	116	4	ABG24038 Mycobacte
35	45	42.5	287	7	ADB80057 Human gua
36	45	42.5	334	4	ABU11182 Human pol
37	45	42.5	334	4	AAy79690 Human pol
38	45	42.5	403	4	AAy40847 Human pol
39	45	42.5	456	4	AAy78706 Human pro
40	44	41.5	506	4	AAy39061 Human pol
41	44	41.5	565	4	ABU71810 Drosophil
42	44	41.5	579	4	AAy76817 Corynebac
43	44	41.5	940	4	AAy90917 C glutami
44	44	41.5	968	4	AAU28194 Novel hum
45	44	41.5	1663	6	AAU28382 Novel hum
					ABO53095 Human put

#### ALIGNMENTS

##### RESULT 1

AAy84265

ID AAY84265 standard; peptide; 20 AA.

XX AAY84265;

XX AAY84265;

DT 12-JUL-2000 (first entry)

XX

DE Truncated macrophage colony stimulating factor tumour antigen.

XX tumour rejection antigen; macrophage colony stimulating gene;

KW macrophage-colony stimulating factor; antigen presenting cell;

KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Homo sapiens.

OS WO200013699-A1.

XX

PD 16-MAR-2000.

XX

PF 03-SEP-1999; 99WO-US020344.

XX

PR 04-SEP-1998; 98US-0099077P.

XX

PA (LUDW-) LUDWIG INST CANCER RES.

XX

PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX

DR WPI; 2000-256859/22.

XX

DR N-PSDB; AA299675.

XX

PT Isolated polypeptide used to treat subjects having a disorder

PT characterized by expression of alternative open reading frame macrophage-

XX colony stimulating factor comprises 25 amino acid residue sequence.

XX

PS Claim 3; Page 64; 74pp; English.

XX

The present sequence represents a truncated tumour rejection antigen precursor, and is encoded by a truncated alternative open reading frame (ORF) of human macrophage colony stimulating gene. Peptides derived from the alternative ORF of macrophage-colony stimulating factor, when presented by an antigen presenting cell having a human leukocyte antigen (HLA) class I molecule, effectively induce the activation and proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF of macrophage-colony stimulating factor are useful for enriching selectively a population of T lymphocytes with CD8+ T lymphocytes. They are also useful for diagnosing a disorder characterized by expression of the polypeptide, and for identifying

CC functional variants and mimetics

```

XX
SQ Sequence 20 AA;
    Query Match      100.0%; Score 106; DB 3; Length 20;
    Best Local Similarity 100.0%; Pred. No. 8.7e-09;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAGLPAVVGSLSPGQEQYHRG 20
Db 1 MAGLPAVVGSLSPGQEQYHRG 20
|||||

RESULT 2
ID AAY84264
AC AAY84264;
XX
XX 12-JUL-2000 (first entry)
XX
XX Peptide of alternate reading frame of macrophage colony stimulating gene.
XX
XX Renal cell carcinoma; antigen; cytotoxic T lymphocyte;
XX tumour rejection antigen; macrophage colony stimulating gene;
XX macrophage-colony stimulating factor; antigen presenting cell;
XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.
XX
XX Homo sapiens.
XX
XX WO200013699-A1.
XX
XX 16-MAR-2000.
XX
XX 03-SEP-1999; 99WO-US020344.
XX
XX 04-SEP-1998; 98US-0099077P.
XX
XX (LUDW-) LUDWIG INST CANCER RES.
XX
XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;
XX
XX WPI; 2000-256859/22.
XX
XX Isolated polypeptide used to treat subjects having a disorder
XX characterized by expression of alternative open reading frame macrophage-
XX colony stimulating factor comprises 25 amino acid residue sequence.
XX
XX Example 1; Page 64; 74pp; English.
XX
XX The present sequence represents a tumour rejection antigen precursor, and
XX is encoded by an alternative open reading frame (ORF) of human macrophage
XX colony stimulating gene. Peptides derived from the alternative ORF of
XX macrophage-colony stimulating factor, when presented by an antigen
XX presenting cell having a human leukocyte antigen (HLA) class I molecule,
XX effectively induce the activation and proliferation of CD8+ cytotoxic T
XX lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF
XX of macrophage-colony stimulating factor are useful for enriching
XX selectively a population of T lymphocytes with CD8+ T lymphocytes. They
XX are also useful for diagnosing a disorder characterized by expression of
XX the polypeptide, and for identifying functional variants and mimetics
XX
XX Sequence 25 AA;
    Query Match      100.0%; Score 106; DB 3; Length 25;
    Best Local Similarity 100.0%; Pred. No. 1.1e-08;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAGLPAVVGSLSPGQEQYHRG 20
Db 1 MAGLPAVVGSLSPGQEQYHRG 20
|||||

CC functional variants and mimetics
XX
SQ Sequence 20 AA;
    Query Match      100.0%; Score 106; DB 3; Length 20;
    Best Local Similarity 100.0%; Pred. No. 8.7e-09;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAGLPAVVGSLSPGQEQYHRG 20
Db 1 MAGLPAVVGSLSPGQEQYHRG 20
|||||

RESULT 3
ID AAY84269
AC AAY84269;
XX
XX 12-JUL-2000 (first entry)
XX
XX Peptide derived from macrophage colony stimulating gene alternative ORF.
XX tumour rejection antigen; macrophage colony stimulating gene;
XX macrophage-colony stimulating factor; antigen presenting cell;
XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.
XX
XX Synthetic.
XX
XX Homo sapiens.
XX
XX WO200013699-A1.
XX
XX 16-MAR-2000.
XX
XX 03-SEP-1999; 99WO-US020344.
XX
XX 04-SEP-1998; 98US-0099077P.
XX
XX (LUDW-) LUDWIG INST CANCER RES.
XX
XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;
XX
XX WPI; 2000-256859/22.
XX
XX Isolated polypeptide used to treat subjects having a disorder
XX characterized by expression of alternative open reading frame macrophage-
XX colony stimulating factor comprises 25 amino acid residue sequence.
XX
XX Example 2; Page 40; 74pp; English.
XX
XX The present sequence represents a peptide which is derived from a tumour
XX rejection antigen precursor encoded by an alternative open reading frame
XX (ORF) of human macrophage colony stimulating gene. Peptides derived from
XX the alternative ORF of macrophage-colony stimulating factor, when
XX presented by an antigen presenting cell having a human leukocyte antigen
XX (HLA) class I molecule, effectively induce the activation and
XX proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic
XX acids derived from the alternate ORF of macrophage-colony stimulating
XX factor are useful for enriching selectively a population of T lymphocytes
XX with CD8+ T lymphocytes. They are also useful for diagnosing a disorder
XX characterized by expression of the polypeptide, and for identifying
XX functional variants and mimetics
XX
XX Sequence 15 AA;
    Query Match      70.8%; Score 75; DB 3; Length 15;
    Best Local Similarity 100.0%; Pred. No. 0.00026;
    Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AGLPAVVGSLSPGQEQE 16
Db 1 AGLPAVVGSLSPGQEQE 15
|||||

RESULT 4
ID AAY84266
AC AAY84266;
XX
XX 12-JUL-2000 (first entry)
XX
XX Peptide derived from macrophage colony stimulating gene alternative ORF.
XX tumour rejection antigen; macrophage colony stimulating gene;

```

KW macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 OS Synthetic.  
 XX Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 XX 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 XX Claim 2; Page 39; 74pp; English.  
 XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 14 AA;  
 XX Query Match 67.9%; Score 72; DB 3; Length 14;  
 XX Best Local Similarity 100.0%; Pred. No. 0.00069;  
 XX Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 4 LPAAVGLSPGQEY 17  
 Db 1 LPAAVGLSPGQEY 14  
 RESULT 5  
 AAY84267  
 ID AAY84267 standard; peptide; 13 AA.  
 AC AAY84267;  
 XX 12-JUL-2000 (first entry)  
 XX Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX tumour rejection antigen; macrophage colony stimulating gene;  
 XX macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 XX Synthetic.  
 XX Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 XX 03-SEP-1999; 99WO-US020344.  
 XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 XX Claim 2; Page 39; 74pp; English.  
 XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 XX Query Match 64.2%; Score 68; DB 3; Length 13;  
 XX Best Local Similarity 100.0%; Pred. No. 0.0025;  
 XX Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 5 PAAVGLSPGQEY 17  
 Db 1 PAAVGLSPGQEY 13  
 RESULT 6  
 AAY84268  
 ID AAY84268 standard; peptide; 13 AA.  
 AC AAY84268;  
 XX 12-JUL-2000 (first entry)  
 XX Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX tumour rejection antigen; macrophage colony stimulating gene;  
 XX macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 XX Synthetic.  
 XX Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 XX 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 XX Claim 2; Page 40; 74pp; English.  
 XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 XX Query Match 64.2%; Score 68; DB 3; Length 13;  
 XX Best Local Similarity 100.0%; Pred. No. 0.0025;  
 XX Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 5 PAAVGLSPGQEY 17  
 Db 1 PAAVGLSPGQEY 13  
 RESULT 6  
 AAY84268  
 ID AAY84268 standard; peptide; 13 AA.  
 AC AAY84268;  
 XX 12-JUL-2000 (first entry)  
 XX Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX tumour rejection antigen; macrophage colony stimulating gene;  
 XX macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 XX Synthetic.  
 XX Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 XX 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 XX Claim 2; Page 40; 74pp; English.  
 XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 XX Query Match 64.2%; Score 68; DB 3; Length 13;  
 XX Best Local Similarity 100.0%; Pred. No. 0.0025;  
 XX Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 5 PAAVGLSPGQEY 17  
 Db 1 PAAVGLSPGQEY 13  
 RESULT 6  
 AAY84268  
 ID AAY84268 standard; peptide; 13 AA.  
 AC AAY84268;  
 XX 12-JUL-2000 (first entry)  
 XX Peptide derived from macrophage colony stimulating gene alternative ORF.  
 XX tumour rejection antigen; macrophage colony stimulating gene;  
 XX macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 XX Synthetic.  
 XX Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 XX 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour

CC rejection antigen precursor encoded by an alternative open reading frame

CC (ORF) of human macrophage colony stimulating gene. Peptides derived from

CC the alternative ORF of macrophage-colony stimulating factor, when

CC presented by an antigen presenting cell having a human leukocyte antigen

CC (HLA) class I molecule, effectively induce the activation and

CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic

CC acids derived from the alternate ORF of macrophage-colony stimulating

CC factor are useful for enriching selectively a population of T lymphocytes

CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder

CC characterized by expression of the polypeptide, and for identifying

CC functional variants and mimetics

XX

SQ Sequence 13 AA;

Query Match 61.3%; Score 65; DB 3; Length 13;

Best Local Similarity 100.0%; Pred. No. 0.007;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 LPAVVGLSPGEQEQ 16

DB 1 LPAVVGLSPGEQEQ 13

|||||

|||||

RESULT 7

ABU49441

ID ABU49441 standard; protein; 277 AA.

XX

AC ABU49441;

DT 19-JUN-2003 (first entry)

XX

DE Protein encoded by Prokaryotic essential gene #34968.

XX

XX Antisense; prokaryotic essential gene; cell proliferation; drug design.

KW

XX

XX Vibrio cholerae.

OS

XX

XX WO200277183-A2.

FN

XX

XX 03-OCT-2002.

PD

XX

XX 21-MAR-2002; 2002WO-US009107.

PF

XX

XX 21-MAR-2001; 2001US-00815242.

PR

XX

XX 06-SEP-2001; 2001US-00948993.

PR

XX

XX 25-OCT-2001; 2001US-0342923P.

PR

XX

XX 08-FEB-2002; 2002US-00072851.

PR

XX

XX 06-MAR-2002; 2002US-0362699P.

PR

XX

XX (ELIT-) ELITRA PHARM INC.

PA

XX

XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;

PI

XX

XX Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

PI

XX

XX WPI; 2003-029926/02.

DR

XX

XX N-PSDB; ACA53311.

DR

XX

XX New antisense nucleic acids, useful for identifying proteins or screening

PT for homologous nucleic acids required for cellular proliferation to

PT isolate candidate molecules for rational drug discovery programs.

PT

XX

XX Claim 25; SEQ ID NO 77365; 1766pp; English.

PS

XX

XX The invention relates to an isolated nucleic acid comprising any one of

CC the 6213 antisense sequences given in the specification where expression

CC of the nucleic acid inhibits proliferation of a cell. Also included are:

CC (1) a vector comprising a promoter operably linked to the nucleic acid

CC encoding a polypeptide whose expression is inhibited by the antisense

CC nucleic acid; (2) a host cell containing the vector; (3) an isolated

CC polypeptide or its fragment whose expression is inhibited by the

CC

CC antisense nucleic acid; (4) an antibody capable of specifically binding

CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular

CC proliferation or the activity of a gene in an operon required for

CC proliferation; (7) identifying a compound that influences the activity of

CC the gene product or that has an activity against a biological pathway;

CC required for proliferation, or that inhibits cellular proliferation; (8)

CC identifying a gene required for cellular proliferation or the biological

CC pathway in which a proliferation-required gene or its gene product lies

CC or a gene on which the test compound that inhibits proliferation of an

CC organism acts; (9) manufacturing an antibiotic; (10) profiling a

CC compound's activity; (11) a culture comprising strains in which the gene

CC product is overexpressed or underexpressed; (12) determining the extent

CC to which each of the strains is present in a culture or collection of

CC strains; or (13) identifying the target of a compound that inhibits the

CC proliferation of an organism. The antisense nucleic acids are useful for

CC identifying proteins or screening for homologous nucleic acids required

CC for cellular proliferation to isolate candidate molecules for rational

CC drug discovery programs, or for screening homologous nucleic acids

CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,

CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is encoded by one of

CC the target prokaryotic essential genes. Note: The sequence data for this

CC patent did not form part of the printed specification, but was obtained

CC in electronic format directly from WIPO at

CC ftp.wipo.int/pub/published\_pot\_sequences

XX

SQ Sequence 277 AA;

Query Match 47.2%; Score 50; DB 6; Length 277;

Best Local Similarity 52.9%; Pred. No. 33;

Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

QY 4 LPAVVGLSPGEQEQYHRG 20

DB 190 LPAFIGAKPYQQAYERG 206

|||||

|||||

RESULT 8

AAB69296

ID AAB69296 standard; protein; 191 AA.

XX

AC AAB69296;

XX

XX 12-SEP-2003 (revised)

DT

XX

XX 20-APR-2001 (first entry)

DT

XX

XX HIV-1 non-subtype B clone 94CY032-3 vif protein.

DE

XX

XX HIV-1; human immunodeficiency virus; non-subtype B; gag; pol; env; vpr;

KW

XX

XX vif; vpr; tat; rev; nef; vaccine.

KW

XX

XX Human immunodeficiency virus 1.

OS

XX

XX WO200026416-A1.

PN

XX

XX 11-MAY-2000.

PD

XX

XX 25-OCT-1999; 99WO-US024837.

PF

XX

XX 02-NOV-1998; 98US-00184418.

PR

XX

XX (UABR-) UAB RES FOUND.

PA

XX

XX Hahn BH, Shaw GM, Gao F;

PI

XX

XX WPI; 2000-365651/31.

DR

XX

XX Novel genomic nucleic acids of non-subtype B human immunodeficiency virus

PT type 1 useful for detecting and treating AIDS comprises a specific

PT nucleotide sequence.

PT

XX

XX Claim 41; Fig 16; 131pp; English.

PS

XX

XX The present in invention provides the protein and coding sequences for a

```
CC number of human immunodeficiency virus (HIV) type 1 non-subtype B
CC isolates. The sequences shown include the near full-length coding
CC sequences from each isolate, and the env, pol, vif, vpr, vpu, gag, tat,
CC rev and nef proteins. These can be used to detect the presence of HIV-1
CC in a sample and to produce antibodies against non-subtype B HIV-1 virus.
CC These antibodies can be used in vaccines to prevent and treat HIV
CC infection. (Updated on 12-SEP-2003 to standardise OS field)
XX
SQ Sequence 191 AA;
    Query Match          46.2%; Score 49; DB 3; Length 191;
    Best Local Similarity 66.7%; Pred. No. 31;
    Matches 8; Conservative 2; Mismatches 0; Gaps 0;
    QY 9 GLSPGEQYHRG 20
    Db 70 GLQPGEDWHLG 81
    |||||:::|
    |||||:::|

RESULT 9
ID ABU30626 standard; protein; 278 AA.
XX
AC ABU30626;
XX
DT 19-JUN-2003 (first entry)
XX
DE Protein encoded by Prokaryotic essential gene #16153.
XX
KW Antisense; prokaryotic essential gene; cell proliferation; drug design.
XX
OS Haemophilus influenzae.
XX
FN WO200277183-A2.
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
XX
PR 06-SEP-2001; 2001US-00948993.
XX
PR 25-OCT-2001; 2001US-0342923P.
XX
PR 08-FEB-2002; 2002US-00072851.
XX
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
XX
PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
XX
DR N-PSDB; ACA34496.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 25; SEQ ID NO 58550; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
```

```
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than S. aureus, S. typhimurium,
CC K. pneumoniae or P. aeruginosa. The present sequence is encoded by one of
CC the target prokaryotic essential genes. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 278 AA;
    Query Match          46.2%; Score 49; DB 6; Length 278;
    Best Local Similarity 52.9%; Pred. No. 46;
    Matches 9; Conservative 2; Mismatches 6; Indels 0; Gaps 0;
    QY 4 LPVVGLSPGEQYHRG 20
    Db 191 LPAPIGAKPYQAYVRG 207
    |||||:::|
    |||||:::|

RESULT 10
ID ABU39195 standard; protein; 278 AA.
XX
AC ABU39195;
XX
DT 19-JUN-2003 (first entry)
XX
DE Protein encoded by Prokaryotic essential gene #24722.
XX
KW Antisense; prokaryotic essential gene; cell proliferation; drug design.
XX
OS Pasteurella multocida.
XX
FN WO200277183-A2.
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
XX
PR 06-SEP-2001; 2001US-00948993.
XX
PR 25-OCT-2001; 2001US-0342923P.
XX
PR 08-FEB-2002; 2002US-00072851.
XX
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
XX
PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
XX
DR N-PSDB; ACA43065.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids, required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 25; SEQ ID NO 67119; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
```

[illegible]

```
Query Match          45.3%; Score 48; DB 6; Length 287;
Best Local Similarity 52.9%; Pred. No. 68;
Matches      9; Conservative    3; Mismatches     0; Gaps      0;
```

---

```
QY           4 LPAVVGLSPGCEQYEYHRG 20
              ||||| : | : | : |
Db            200 LPAFVGANPYKQAYEKG 216
```

---

```
RESULT 12
ADA36829
ID ADA36829 standard; protein; 303 AA.
XX XX AC AC AC AC
DT DT 20-NOV-2003 (first entry)
XX XX Acinetobacter baumannii protein #3990.
XX DE Acinetobacter baumannii; bacterial disease; antibacterial; vaccine;
KW KW plant biocontrol agent.
XX OS Acinetobacter baumannii.
XX PN US6562958-B1.
XX PD 13-MAY-2003.
XX PF 04-JUN-1999; 99US-00328352.
XX PR 09-JUN-1998; 98US-0088701P.
XX PA (GENO-) GENOME THERAPEUTICS CORP.
```

XX Breton G, Bush D;  
 XX WPI; 2003-576092/54.  
 DR N-PSDB; ADA32703.  
 XX New Acinetobacter baumannii proteins and nucleic acids, useful as reagents  
 PT for diagnosing a bacterial disease, as components of antibacterial  
 PT vaccines, as targets for antibacterial drugs, or as biocontrol agents for  
 PT plants.  
 PS Example; SEQ ID NO 8116; 328pp; English.  
 XX The invention relates to isolated Acinetobacter baumannii nucleic acids.  
 CC The A. baumannii nucleic acids and polypeptides are useful as reagents  
 CC for diagnosing a bacterial disease, as components of antibacterial  
 CC vaccines, as targets for antibacterial drugs, to detect the presence of  
 CC A. baumannii and other Acinetobacter species in a sample, in screening  
 CC compounds for the ability to interfere with the A. baumannii life cycle  
 CC or to inhibit A. baumannii infection, and as biocontrol agents for  
 CC plants. The present sequence represents the amino acid sequence of an A.  
 CC baumannii protein.  
 XX Sequence 303 AA;  
 SQ

Query Match 45.3%; Score 48; DB 6; Length 303;  
 Best Local Similarity 52.9%; Pred. No. 72;  
 Matches 9; Conservative 3; Mismatches 5; Indels 0; Gaps 0;  
 QY 4 LPAVGLSPGEQYHRG 20  
 DB 216 LPAFGANPYQAYEKG 232

RESULT 13  
 ABUS0765  
 ID ABUS0765 standard; protein; 151 AA.  
 XX  
 AC ABUS0765;  
 XX  
 DT 07-MAY-2003 (first entry)  
 XX Helicobacter pylori selected interacting domain (SID) protein #108.  
 DE Protein-protein interaction; ulcer; selected interacting domain; SID.  
 XX Helicobacter pylori.  
 OS Helicobacter pylori.  
 XX WO200266501-A2.  
 XX 29-AUG-2002.  
 XX 28-DEC-2001; 2001WO-EP015428.  
 XX 02-JAN-2001; 2001US-0259302P.  
 XX (HYBR-) HYBRIGENICS.  
 PA (INSP ) INST PASTEUR.  
 XX Legrain P, Rain J, Colland F, De Reuse H, Labigne A;  
 WPI; 2002-674910/72.  
 N-PSDB; ABX65509.  
 XX New complexes of protein-protein interactions in Helicobacter pylori,  
 PT useful for identifying modulating compounds for treating or preventing  
 PT ulcers in mammals.  
 XX Claim 6; Page 111; 642pp; English.  
 XX The invention describes a complex of protein-protein interactions in  
 CC Helicobacter pylori selected from 421 complexes given in the  
 CC specification. The complex of protein-protein interactions are useful for  
 CC screening for agents which modulate the interaction of proteins.  
 CC Modulating compounds which binds to a targeted bacterial protein may be  
 CC used for treating or preventing ulcers in a human or animal. This is the  
 CC amino acid sequence of a selected interacting domain (SID), identified  
 CC via protein-protein interactions. Note: Where the patent number printed  
 CC at the top of the pages in the specification has obscured areas of  
 CC protein sequence, the indexer has replaced the residue with an X to  
 CC represent an illegible residue  
 XX Sequence 151 AA;  
 SQ

Query Match 44.3%; Score 47; DB 5; Length 151;  
 Best Local Similarity 47.1%; Pred. No. 48;  
 Matches 8; Conservative 4; Mismatches 5; Indels 0; Gaps 0;  
 QY 4 LPAVGLSPGEQYHRG 20  
 DB 79 LPAFGANPYQAFERG 95

RESULT 14  
 ABUS1750  
 ID ABUS1750 standard; protein; 155 AA.  
 XX  
 AC ABUS1750;  
 XX  
 DT 07-MAY-2003 (first entry)  
 XX Helicobacter pylori selected interacting domain (SID) protein #1094.  
 DE Protein-protein interaction; ulcer; selected interacting domain; SID.  
 XX Helicobacter pylori.  
 OS Helicobacter pylori.  
 XX WO200266501-A2.  
 XX 29-AUG-2002.  
 XX 28-DEC-2001; 2001WO-EP015428.  
 XX 02-JAN-2001; 2001US-0259302P.  
 XX (HYBR-) HYBRIGENICS.  
 PA (INSP ) INST PASTEUR.  
 XX Legrain P, Rain J, Colland F, De Reuse H, Labigne A;  
 WPI; 2002-674910/72.  
 N-PSDB; ABX66495.  
 XX New complexes of protein-protein interactions in Helicobacter pylori,  
 PT useful for identifying modulating compounds for treating or preventing  
 PT ulcers in mammals.  
 XX Claim 6; Page 354; 642pp; English.  
 XX The invention describes a complex of protein-protein interactions in  
 CC Helicobacter pylori selected from 421 complexes given in the  
 CC specification. The complex of protein-protein interactions are useful for  
 CC screening for agents which modulate the interaction of proteins.  
 CC Modulating compounds which binds to a targeted bacterial protein may be  
 CC used for treating or preventing ulcers in a human or animal. This is the  
 CC amino acid sequence of a selected interacting domain (SID), identified  
 CC via protein-protein interactions. Note: Where the patent number printed  
 CC at the top of the pages in the specification has obscured areas of  
 CC protein sequence, the indexer has replaced the residue with an X to  
 CC represent an illegible residue  
 XX Sequence 155 AA;  
 SQ

Query Match 44.3%; Score 47; DB 5; Length 155;  
 Best Local Similarity 47.1%; Pred. No. 49;  
 Matches 8; Conservative 4; Mismatches 5; Indels 0; Gaps 0;  
 QY 4 LPAVGLSPGEQYHRG 20  
 DB 79 LPAFGANPYQAFERG 95

RESULT 15  
 ABUS1750  
 ID ABUS1750 standard; protein; 155 AA.  
 XX  
 AC ABUS1750;  
 XX  
 DT 07-MAY-2003 (first entry)  
 XX Helicobacter pylori selected interacting domain (SID) protein #1094.  
 DE Protein-protein interaction; ulcer; selected interacting domain; SID.  
 XX Helicobacter pylori.  
 OS Helicobacter pylori.  
 XX WO200266501-A2.  
 XX 29-AUG-2002.  
 XX 28-DEC-2001; 2001WO-EP015428.  
 XX 02-JAN-2001; 2001US-0259302P.  
 XX (HYBR-) HYBRIGENICS.  
 PA (INSP ) INST PASTEUR.  
 XX Legrain P, Rain J, Colland F, De Reuse H, Labigne A;  
 WPI; 2002-674910/72.  
 N-PSDB; ABX66495.  
 XX New complexes of protein-protein interactions in Helicobacter pylori,  
 PT useful for identifying modulating compounds for treating or preventing  
 PT ulcers in mammals.  
 XX Claim 6; Page 354; 642pp; English.  
 XX The invention describes a complex of protein-protein interactions in  
 CC Helicobacter pylori selected from 421 complexes given in the  
 CC specification. The complex of protein-protein interactions are useful for  
 CC screening for agents which modulate the interaction of proteins.  
 CC Modulating compounds which binds to a targeted bacterial protein may be  
 CC used for treating or preventing ulcers in a human or animal. This is the  
 CC amino acid sequence of a selected interacting domain (SID), identified  
 CC via protein-protein interactions. Note: Where the patent number printed  
 CC at the top of the pages in the specification has obscured areas of  
 CC protein sequence, the indexer has replaced the residue with an X to  
 CC represent an illegible residue  
 XX Sequence 155 AA;  
 SQ

Query Match 44.3%; Score 47; DB 5; Length 155;  
 Best Local Similarity 47.1%; Pred. No. 49;  
 Matches 8; Conservative 4; Mismatches 5; Indels 0; Gaps 0;  
 QY 4 LPAVGLSPGEQYHRG 20  
 DB 79 LPAFGANPYQAFERG 95

RESULT 16  
 ABUS1750  
 ID ABUS1750 standard; protein; 155 AA.  
 XX  
 AC ABUS1750;  
 XX  
 DT 07-MAY-2003 (first entry)  
 XX Helicobacter pylori selected interacting domain (SID) protein #1094.  
 DE Protein-protein interaction; ulcer; selected interacting domain; SID.  
 XX Helicobacter pylori.  
 OS Helicobacter pylori.  
 XX WO200266501-A2.  
 XX 29-AUG-2002.  
 XX 28-DEC-2001; 2001WO-EP015428.  
 XX 02-JAN-2001; 2001US-0259302P.  
 XX (HYBR-) HYBRIGENICS.  
 PA (INSP ) INST PASTEUR.  
 XX Legrain P, Rain J, Colland F, De Reuse H, Labigne A;  
 WPI; 2002-674910/72.  
 N-PSDB; ABX66495.  
 XX New complexes of protein-protein interactions in Helicobacter pylori,  
 PT useful for identifying modulating compounds for treating or preventing  
 PT ulcers in mammals.  
 XX Claim 6; Page 354; 642pp; English.  
 XX The invention describes a complex of protein-protein interactions in  
 CC Helicobacter pylori selected from 421 complexes given in the  
 CC specification. The complex of protein-protein interactions are useful for  
 CC screening for agents which modulate the interaction of proteins.  
 CC Modulating compounds which binds to a targeted bacterial protein may be  
 CC used for treating or preventing ulcers in a human or animal. This is the  
 CC amino acid sequence of a selected interacting domain (SID), identified  
 CC via protein-protein interactions. Note: Where the patent number printed  
 CC at the top of the pages in the specification has obscured areas of  
 CC protein sequence, the indexer has replaced the residue with an X to  
 CC represent an illegible residue  
 XX Sequence 155 AA;  
 SQ



QY 4 LPAVGLSPGEQYHRG 20  
Db |||:|:|:|:|:|:|  
80 LPAFIGANPYQQAIFRG 96

RESULT 15  
AAB69298  
ID AAB69298 standard; protein; 192 AA.  
XX AC AAB69298;  
XX 12-SEP-2003 (revised)  
DT 20-APR-2001 (first entry)  
XX DE HIV-1 non-subtype B clone 962M751-3 vif protein.  
XX HIV-1; human immunodeficiency virus; non-subtype B; gag; pol; env; vpr;  
KW vif; vpr; tat; rev; nef; vaccine.  
XX OS Human immunodeficiency virus 1.  
XX WO200026416-A1.  
PN 11-MAY-2000.  
XX 25-OCT-1999; 99WO-US024837.  
XX 02-NOV-1998; 98US-00184418.  
XX (UABR-) UAB RES FOUND.  
XX Hahn BH, Shaw GM, Gao F;  
PI WPI; 2000-365651/31.  
XX Novel genomic nucleic acids of non-subtype B human immunodeficiency virus  
PT type 1 useful for detecting and treating AIDS comprises a specific  
PT nucleotide sequence.  
XX Claim 41; Fig 16; 131pp; English.  
XX The present invention provides the protein and coding sequences for a  
CC number of human immunodeficiency virus (HIV) type 1 non-subtype B  
CC isolates. The sequences shown include the near full-length coding  
CC sequences from each isolate, and the env, pol, vif, vpr, vpu, gag, tat,  
CC rev and nef proteins. These can be used to detect the presence of HIV-1  
CC in a sample and to produce antibodies against non-subtype B HIV-1 virus.  
CC These antibodies can be used in vaccines to prevent and treat HIV  
CC infection. (Updated on 12-SEP-2003 to standardise OS field)  
XX SQ Sequence 192 AA;  
Query Match 44.3%; Score 47; DB 3; Length 192;  
Best Local Similarity 66.7%; Pred. NO. 62;  
Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
QY 9 GLSPGEQYHRG 20  
Db |||:|:|:|:|:|:|  
71 GLHFGEREWHUG 82

Search completed: May 7, 2004, 12:33:43  
Job time : 56.8 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:27:20 ; Search time 8.96 Seconds  
(without alignments)  
150.299 Million cell updates/sec

Title: US-09-786-214A-12  
Perfect score: 72  
Sequence: 1 LPAVVGLSPGRQY 14

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 283366 segs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : PIR 78: \*  
1: pir1: \*  
2: pir2: \*  
3: pir3: \*  
4: pir4: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	44	61.1	381	2 F75270	cytochrome P450 -
2	43	59.7	211	2 A64490	precorrin-6Y methy
3	43	59.7	623	2 T40391	probable lysophosp
4	43	59.7	821	2 C84304	DNA helicase limpo
5	41	56.9	156	2 F87551	conserved hypochet
6	41	56.9	326	2 T45226	probable N5,N10-me
7	41	56.9	508	2 E70764	probable cobi prot
8	40	55.6	227	2 B90400	hypothetical prote
9	40	55.6	243	2 I54459	MEC H-2K1-k - mous
10	40	55.6	428	2 AG1304	uracil permease ho
11	40	55.6	428	2 AG1676	uracil permease ho
12	40	55.6	540	2 A75250	carboxylesterase,
13	40	55.6	661	2 G84511	hypothetical prote
14	40	55.6	673	2 T50281	probable lysophosp
15	39	54.2	43	2 S21065	Ig kappa chain v r
16	39	54.2	96	2 S45441	Ig kappa chain v r
17	39	54.2	103	2 S19975	Ig kappa chain v r
18	39	54.2	106	2 PS0070	Ig kappa chain v r
19	39	54.2	106	2 PC4282	Ig kappa chain (an
20	39	54.2	107	2 S57444	Ig kappa chain v r
21	39	54.2	108	2 C30502	Ig kappa chain v r
22	39	54.2	108	2 S39988	Ig kappa chain v r
23	39	54.2	108	2 G44151	Ig kappa chain v r
24	39	54.2	111	2 S23628	Ig kappa chain v r
25	39	54.2	114	2 S54905	Ig kappa chain v r
26	39	54.2	115	1 K3HUVG	Ig kappa chain pre
27	39	54.2	115	1 KYMSL7	Ig kappa chain pre
28	39	54.2	115	2 S11697	Ig kappa chain pre
29	39	54.2	116	2 B25521	Ig kappa chain pre

30 39 54.2 119 2 S41816  
31 39 54.2 125 2 S40344  
32 39 54.2 128 2 PNO445  
33 39 54.2 128 2 S40379  
34 39 54.2 128 2 A56701  
35 39 54.2 129 2 S29627  
36 39 54.2 129 2 S40363  
37 39 54.2 132 2 S05288  
38 39 54.2 144 2 P0106  
39 39 54.2 144 2 B30502  
40 39 54.2 215 2 A23746  
41 39 54.2 277 2 P82130  
42 39 54.2 278 2 P64131  
43 39 54.2 319 2 AD0941  
44 39 54.2 563 2 JQ0623  
45 39 54.2 1367 2 S74285

## ALIGNMENTS

## RESULT 1

F75270

Cytochrome P450 - Deinococcus radiodurans (strain R1)

C:Species: Deinococcus radiodurans  
C:Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000

C:Accession: F75270  
R:White, O.; Eissen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;  
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.; M;  
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.

Science 286, 1571-1577, 1999

A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.

A:Reference number: A75250; MUID:20036896; PMID:10567266

A:Accession: F75270

A>Status: Preliminary

A:Molecule type: DNA

A:Residues: 1-391 <WHI>

A:Cross-references: GB:AE002076; GB:AE000513; NID:g6460285; PIDN:AAF12016.1; PID:g64602;

A:Experimental source: strain R1

C:Genetics:

A:Gene: DR2473

A:Map position: 1

Query Match 61.1%; Score 44; DB 2; Length 381;  
Best Local Similarity 100.0%; Pred. No. 12;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGLSP 9

Db 52 LPAVVGLSP 60

## RESULT 2

A64490

precorrin-6Y methylase homolog - Methanococcus jannaschii

C:Species: Methanococcus jannaschii

C:Date: 13-Sep-1996 #sequence\_revision 13-Sep-1996 #text\_change 18-Aug-2003

C:Accession: A64490

R:Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann, R.D.; Sutton, G.G.; Blake,  
; Reich, C.I.; Overbeek, R.; Kirkness, E.F.; Weinstock, K.G.; Merrick, J.M.; Glodek, A.  
rson, J.D.; Sadow, P.W.; Hanna, M.C.; Cotton, M.D.; Roberts, K.M.; Hurst, M.A.

Science 273, 1058-1073, 1996

A:Authors: Kaine, B.P.; Borodovsky, M.; Klenk, H.P.; Fraser, C.M.; Smith, H.O.; Woese, C.

A:Title: Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii.

A:Reference number: A64300; MUID:96337999; PMID:8688087

A:Accession: A64490

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-211 <BUL>

A:Cross-references: GB:U67593; GB:L77117; NID:g2826427; PIDN:AAB99541.1; PID:g1592152;

C:Genetics:

A:Map position: FOR1500322-1500957

C:Superfamily: precorrin-6Y methylase CbiE

Query Match 59.7%; Score 43; DB 2; Length 211;  
 Best Local Similarity 54.5%; Pred. No. 9.3;  
 Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGGEQY 14  
 :||: |||:  
 Db 4 IVGIGPGDREY 14

RESULT 3  
 T40991  
 probable lysophospholipase precursor - fission yeast (*Schizosaccharomyces pombe*)  
 C:Species: *Schizosaccharomyces pombe*  
 C>Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 21-Jan-2000  
 C:Accession: T40991  
 R:Lyne, M.; Rajandream, M.A.; Barrell, B.G.; Volckaert, G.  
 submitted to the EMBL Data Library, March 1999  
 A:Reference number: Z21962  
 A:Accession: T40991  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-623 <LYN>  
 A:Cross-references: EMBL:AL049559; PIDN:CA840176.1; GSPDB:GN000068; SPDB:SPCC1450.09c  
 A:Experimental source: strain 972h; cosmid ci450  
 C:Genetics:  
 A:Gene: SPDB:SPCC1450.09c  
 A:Map position: 3  
 C:Superfamily: yeast lysophospholipase

Query Match 59.7%; Score 43; DB 2; Length 623;  
 Best Local Similarity 69.2%; Pred. No. 29;  
 Matches 9; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 2 PAVGLSPGGEQY 14  
 || ||| |||:  
 Db 76 PASDGLSTGEQRF 88

RESULT 4  
 CB4304  
 DNA helicase [imported] - *Halobacterium* sp. NRC-1  
 C:Species: *Halobacterium* sp. NRC-1  
 C>Date: 02-Feb-2001 #sequence\_revision 02-Feb-2001 #text\_change 02-Feb-2001  
 C:Accession: CB4304  
 R:Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.; Leithausen, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jablo  
 Jung, K.H.; Alam, M.; Freitas, T.  
 Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000  
 A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li  
 A:Title: Genome sequence of *Halobacterium* species NRC-1.  
 A:Reference number: A84160; MUID:20504483; PMID:11016950  
 A:Accession: CB4304  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-821 <STO>  
 A:Cross-references: GB:AE004437; NID:gl0580995; PIDN:AAG19799.1; GSPDB:GN00138  
 C:Genetics:  
 A:Gene: hel

Query Match 59.7%; Score 43; DB 2; Length 821;  
 Best Local Similarity 90.0%; Pred. No. 38;  
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGGEQ 12  
 ||||| |||:  
 Db 326 AVVGLSPAEG 335

RESULT 5  
 F87551  
 conserved hypothetical protein CC2439 [imported] - *Caulobacter crescentus*  
 C:Species: *Caulobacter crescentus*

C>Date: 20-Apr-2001 #sequence\_revision 20-Apr-2001 #text\_change 10-May-2001  
 C:Accession: F87551  
 R:Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.  
 B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolo  
 n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M.  
 Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001  
 A:Title: Complete Genome Sequence of *Caulobacter crescentus*.  
 A:Reference number: A87249; MUID:21173698; PMID:11259647  
 A:Accession: F87551  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-156 <STO>  
 A:Cross-references: GB:AE005673; NID:gl3423984; PIDN:AAK24410.1; GSPDB:GN00148  
 C:Genetics:  
 A:Gene: CC2439  
 C:Superfamily: Haemophilus influenzae conserved hypothetical protein HI0305

Query Match 56.9%; Score 41; DB 2; Length 156;  
 Best Local Similarity 80.0%; Pred. No. 15;  
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGGEQ 12  
 ||||| |||:  
 Db 18 AVVGLDPGEK 27

RESULT 6  
 T45226  
 probable NS<sub>10</sub>-methylene-tetrahydromethanopterin reductase (F420-dependent) [imported]  
 C:Species: *Methanobolus tindarius*  
 C>Date: 21-Jan-2000 #sequence\_revision 21-Jan-2000 #text\_change 21-Jan-2000  
 C:Accession: T45226  
 R:Westenberg, D.J.; Braune, A.; Ruppert, C.; Mueller, V.; Herzberg, C.; Gottschalk, G.;  
 submitted to the EMBL Data Library, September 1998  
 A:Description: The F420H2-dehydrogenase from *Methanobolus tindarius*: Cloning of the ffa  
 A:Reference number: Z22947  
 A:Accession: T45226  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-326 <WES>  
 A:Cross-references: EMBL:AJ011519; PIDN:CA856639.1  
 A:Experimental source: DSM 2278  
 C:Genetics:  
 A:Gene: ffa

Query Match 56.9%; Score 41; DB 2; Length 326;  
 Best Local Similarity 70.0%; Pred. No. 32;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGGEQ 12  
 ||||| |||:  
 Db 88 AILGLPGGEQ 97

RESULT 7  
 E70764  
 probable cobi protein - *Mycobacterium tuberculosis* (strain H37RV)  
 C:Species: *Mycobacterium tuberculosis*  
 C>Date: 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change 22-Oct-1999  
 C:Accession: E70764  
 R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.  
 ; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.  
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.  
 Nature 393, 537-544, 1998  
 A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
 A:Title: Deciphering the biology of *Mycobacterium tuberculosis* from the complete genome  
 A:Reference number: A70500; MUID:98295987; PMID:9634230  
 A:Accession: E70764  
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-508 <COL>  
 A:Cross-references: GB:Z73966; GB:AL123456; NID:g3261577; PIDN:CAA98214.1; PID:e246996;  
 A:Experimental source: strain H37RV

RESULT 10  
AD1304  
uracil permease homolog pyrP [imported] - *Listeria monocytogenes* (strain EGD-e)  
C-Species: *Listeria monocytogenes*

Query Match	55.6%	Score 40;	DB 2;	Length 428;
Best Local Similarity	70.0%	Pred. No. 63;		
Matches	7.	Conservative	2:	Mismatches
				1: Indels
				0: Gaps
				0:

C;Species: *Deinococcus radiodurans*  
C;Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000  
C;Accession: A75250

S.; Smith, H.O.; Venter, J.C.; Fraser, C.M. Science 286, 1571-1577, 1999

A;Reference number: A75250; MUID:20036896; PMID:10567266  
 A;Accession: A75250  
 A;Molecule type: DNA  
 A;Residues: 1-540 <WHI>  
 A;Cross-references: GB:AE002092; GB:AE000513; NID:g6460455; PIDN:AAF12163.1; PID:g646045  
 A;Experimental source: strain R1  
 C;Genetics:

A;Map position: 1  
 A;Gene: DR2626  
 C;Superfamily: cholinesterase; cholinesterase homology

Query Match 55.6%; Score 40; DB 2; Length 540;  
 Best Local Similarity 70.0%; Pred. No. 80;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 PAVVGLSPGCE 11  
 ||:|||||  
 Db 512 PQVGLAPGE 521

## RESULT 13

GB4511

hypothetical protein At2g13900 [imported] - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 02-Feb-2001 #sequence\_revision 02-Feb-2001 #text\_change 02-Feb-2001

C;Accession: G84511

R;Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;  
 M.; Xoo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon, L.  
 eus, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter,  
 Nature 402, 761-768, 1999

A;Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.

A;Reference number: AB4420; MUID:20083487; PMID:10617197

A;Accession: G84511

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-661 <STO>

A;Cross-references: GB:AE002093; NID:g6598598; PIDN:AAF18650.1; GSPDB:GN00139

C;Genetics:

A;Gene: At2g13900

A;Map position: 2

Query Match 55.6%; Score 40; DB 2; Length 661;  
 Best Local Similarity 53.8%; Pred. No. 99;  
 Matches 7; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 PAVVGLSPGCE 14  
 ||:|||||  
 Db 437 PTLTKIVPGCEY 449

## RESULT 14

T50281

probable lysophospholipase (EC 3.1.1.5) precursor SPAC977.09c [similarity] - fission yea

C;Species: Schizosaccharomyces pombe

C;Date: 03-Jun-2000 #sequence\_revision 09-Jun-2000 #text\_change 19-Jan-2001

C;Accession: T50281; T42738

R;Zimmermann, W.; Wambutt, R.; McDougall, R.C.; Rajandream, M.A.; Barrell, B.G.

submitted to the EMBL Data Library, January 2000

A;Reference number: Z25053

A;Accession: T50281

A;Status: translated from GB/EMBL/DDBJ

A;Molecule type: DNA

A;Residues: 1-673 <ZIM>

A;Cross-references: EMBL:AL137130; NID:g6742151; PIDN:CAB69631.1; PID:g6742159; GSPDB:GN

A;Experimental source: strain 972h(-); cosmid c977

R;Yoshioka, S.; Kato, K.; Nakai, K.; Okayama, H.; Nojima, H.

DNA Res. 4, 363-369, 1997

A;Title: Identification of open reading frames in Schizosaccharomyces pombe cDNAs.

A;Reference number: Z17323; MUID:98162722; PMID:9501991

A;Accession: T42738

A;Status: preliminary; translated from GB/EMBL/DDBJ

A;Molecule type: mRNA

A;Residues: 427-673 <YOS>

A;Cross-references: EMBL:D89183; NID:gl749573; PIDN:BAAL3845.1; PID:gl749574  
 A;Experimental source: strain PR745

C;Genetics:

A;Gene: SPDB:SPAC977.09c

A;Map position: 1

A;Introns: 651/3

C;Function:

A;Description: catalyzes the hydrolysis of 2-lysophosphatidylcholine to glycerophosphoch  
 C;Superfamily: yeast lysophospholipase  
 C;Keywords: carboxylic ester hydrolase

Query Match 55.6%; Score 40; DB 2; Length 673;  
 Best Local Similarity 61.5%; Pred. No. 1e+02;  
 Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 2 PAVVGLSPGCE 14  
 ||:|||||  
 Db 83 PASEGLNEGEQSY 95

## RESULT 15

S21065

Ig kappa chain V region (anti-RH(D)) - human (fragment)

C;Species: Homo sapiens (man)

C;Date: 22-Nov-1993 #sequence\_revision 24-May-1996 #text\_change 09-May-1997

C;Accession: S21065

R;Dlouha, A.; Lecroisey, A.; Henschen, A.; Rouger, P.; Keil, B.

Protein Seq. Data Anal. 4, 317-318, 1991

A;Title: Subgroup assignment of a human monoclonal anti-Rh(D) antibody.

A;Reference number: S21065; MUID:92253544; PMID:1812483

A;Accession: S21065

A;Molecule type: protein

A;Residues: 1-43 <DLO>

A;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

Query Match 54.2%; Score 39; DB 2; Length 43;  
 Best Local Similarity 63.6%; Pred. No. 8.4;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGCE 12  
 ||:|||||  
 Db 8 PATLSLSPGER 18

Search completed: May 7, 2004, 12:39:06  
 Job time : 9.12667 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 5.04 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-12

Perfect score: 72

Sequence: 1 LPAVGLSPGEQY 14

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	59.7	211	1 COBL_METJA	Q58917 methanococ
2	43	59.7	633	2 PUB5_SCHPO	Q9y7n6 schizosacch
3	41	56.9	156	1 RUVX_CAUCR	Q9a5k8 caulobacter
4	41	56.9	326	1 MER_METRI	Q9uxp0 methanolobu
5	41	56.9	508	1 COBI_MYCTU	Q10677 mycobacteri
6	40	55.6	673	1 PUB4_SCHPO	Q9p327 schizosacch
7	39	54.2	115	1 KV31_HUMAN	P04433 homo sapien
8	39	54.2	115	1 KV51_MOUSE	P01642 mus musculu
9	39	54.2	278	1 PURU_HAEIN	Q03432 haemophilus
10	39	54.2	516	1 C4AD_DROME	Q9v4t3 drosophila
11	39	54.2	597	1 NR41_RAT	P22829 rattus norv
12	39	54.2	1402	1 N160_MOUSE	Q920W3 mus musculu
13	39	54.2	1636	1 BUD3_YEAST	P25558 saccharomyc
14	38	52.8	390	1 COBL_MYCTU	Q10671 mycobacteri
15	38	52.8	429	1 R51_LEULA	P50889 leuconostoc
16	38	52.8	507	1 CAT4_PICAN	P10263 pichia angu
17	38	52.8	699	1 EFG_HAEIN	P43925 haemophilus
18	38	52.8	700	1 EFG_PASMO	P57938 pasteurella
19	37.5	52.1	827	1 MAK1_MOUSE	P62018 mus musculu
20	37	51.4	280	1 PMXA_MOUSE	Q62066 mus musculu
21	37	51.4	281	1 PMXA_RAT	Q62782 rattus norv
22	37	51.4	446	1 COBJ_ARCFU	Q29534 a cobalanin
23	37	51.4	446	1 ENO1_MAIZE	P26301 zea mays (m
24	37	51.4	557	1 PUR6_VIGAC	P55195 vigna acon
25	37	51.4	607	1 GLMS_CLOTE	Q890u2 c glucosami
26	37	51.4	637	1 MUTL_CAUCR	Q19p66 caulobacter
27	37	51.4	658	1 VGI8_BPT4	P13332 bacterioph
28	37	51.4	753	1 CKXA_BACUF	Q92321 bacillus th
29	37	51.4	813	1 CADM_MOUSE	Q9wt5 mus musculu
30	37	51.4	985	1 4ET_HUMAN	Q9nra8 homo sapien
31	37	51.4	992	1 EVC_HUMAN	P57679 homo sapien
32	37	51.4	2269	1 RRF1_SV41	P35341 simian viru
33	36.5	50.7	374	1 RGSK_BOVIN	P79348 bos taurus

#### RESULT 1

COBL\_METJA STANDARD; PRT; 211 AA.

AC Q58917; 1998 (Rel. 37, Created)  
DT 15-DEC-1998 (Rel. 37, Last sequence update)  
DT 28-FEB-2003 (Rel. 41, Last annotation update)  
DE Probable precorrin-6Y C5,15-methyltransferase [decarboxylating]  
DE (EC 2.1.1.132) (Precorrin-6 methyltransferase) (Precorrin-6Y  
DE methylase)  
DE GN COBL OR M1522.

OS Methanococcus jannaschii.

OC Archaea; Euryarchaeota; Methanococci; Methanococcales;

OC Methanocaldococcaceae; Methanocaldococcus.

OX NCBI\_TaxID=2190;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=JAL-1 / DSM 2661 / ATCC 43067;

RX MEDLINE=96337999; PubMed=8688087;

RA Bult C.G., White O., Olsen G.J., Zhou L., Fleischmann R.D.,

Sutton G.G., Blake J.A., FitzGerald L.M., Clayton R.A., Gocayne J.D.,

Kerlavage A.R., Dougherty B.A., Tomb J.-F., Adams M.D., Reich C.I.,

Overbeek R., Kirkness E.F., Weinstock K.G., Merrick J.M., Glodek A.,

Scott J.L., Geoghegan N.S.M., Weidman J.F., Fuhrmann J.L., Nguyen D.,

Uterback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,

Cotton M.D., Roberts K.M., Hurst M.A., Kaine B.P., Borodovsky M.,

Klenk H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;

"Complete genome sequence of the methanogenic archaeon, Methanococcus

jannaschii".

RL Science 273:1058-1073 (1996).

CC -!- FUNCTION: CATALYZES THE METHYLATION OF BOTH C-5 AND C-15 IN

PRECORRIN-6Y TO FORM PRECORRIN-8X (BY SIMILARITY).

CC -!- CATALYTIC ACTIVITY: 2 S-adenosyl-L-methionine + precorrin-6Y = 2

S-adenosyl-L-homocysteine + precorrin-8X + CO(2).

CC -!- PATHWAY: Cobalamin biosynthesis.

CC -!- SIMILARITY: TO S-TYRPHIMURIUM CBIE; ALSO, LOW, TO OTHER

METHYLASES INVOLVED IN COBALAMIN BIOSYNTHESIS.

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CC EMBL; U67593; AAB93541.1; -.

DR F1R; A64490; A64490.

DR TIGR; M1522; -.

DR InterPro; IPR000878; Cor/por Mettransf.

DR Pfam; PF00590; TP methylase; 1.

DR Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;

KW Methyltransferase; Complete proteome.

SEQUENCE 211 AA; 23805 MW; 279A1A2B14369510 CRC64;

Query Match 59.7%; Score 43; DB 1; Length 211;

Best Local Similarity 54.5%; Pred. No. 5.8;									
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;									
QY	4	VVGLSPGEQY 14							
Db	4	IVGIGPGDREY 14							
RESULT 2									
DR	FLB5_SCHPO	STANDARD;	PRT;	633	AA.				
DR	Q9Y7N6;								
DR	10-OCT-2003 (Rel. 42, Created)								
DR	10-OCT-2003 (Rel. 42, Last sequence update)								
DR	10-OCT-2003 (Rel. 42, Last annotation update)								
DR	Putative lysophospholipase C1450.09c precursor (EC 3.1.1.5)								
DR	(Phospholipase B).								
DR	SPC1450.09c.								
DR	Schizosaccharomyces pombe (Fission yeast).								
DR	Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;								
DR	Schizosaccharomycetales; Schizosaccharomycetaceae;								
DR	Schizosaccharomycetes.								
DR	NCBI_TaxID=4896;								
DR	[1]								
SEQUENCE FROM N.A.									
DR	STRAIN=972;								
DR	MEDLINE=21848401; PubMed=11859360;								
DR	Wood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,								
DR	Sgouros J., Peat N., Hayles J., Baker S., Baeham D., Bowman S.,								
DR	Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,								
DR	Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,								
DR	Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,								
DR	Hollroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,								
DR	James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,								
DR	Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,								
DR	Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,								
DR	Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,								
DR	Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,								
DR	Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,								
DR	Woodward J., Volckaert G., Aert R., Robben J., Grymonprez B.,								
DR	Weltjens I., Vanstreels E., Kieger M., Schaefer M., Mueller-Auer S.,								
DR	Gabel C., Fuchs M., Fritze C., Holzer E., Moestl D., Halbert H.,								
DR	Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,								
DR	Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,								
DR	Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,								
DR	Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,								
DR	Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,								
DR	Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,								
DR	Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,								
DR	Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,								
DR	Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;								
DR	"The genome sequence of Schizosaccharomyces pombe.";								
DR	Nature 415:871-880(2002).								
DR	-1- FUNCTION: Catalyzes the release of fatty acids from								
DR	lysophospholipids (By similarity).								
DR	-1- CATALYTIC ACTIVITY: 2-lysophosphatidylcholine + H(2)O =								
DR	glycerophosphocholine + a fatty acid anion.								
DR	-1- SUBCELLULAR LOCATION: Secreted (Probable).								
DR	-1- SIMILARITY: Belongs to the lysophospholipase family.								
DR	-----								
DR	This SWISS-PROT entry is copyright. It is produced through a collaboration								
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DR	the European Bioinformatics Institute. There are no restrictions on its								
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DR	entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a>								
DR	or send an email to <a href="mailto:license@isb-sib.ch">license@isb-sib.ch</a> ).								
DR	-----								
DR	EMBL; AL049559; CAB40176.2; -								
DR	GeneDB SPombe; SPC1450.09c; -								
DR	InterPro; IPR002642; PLAC.								
DR	Pfam; PF01735; PLA2_B; 1.								
DR	SMART; SM00022; PLAC; 1.								

```

DR EMBL; AB005913; AAK24410.1; -.
DR PIR; F87551; F87551.
DR TIGR; CC2439; -.
DR HAMAP; MF 00651; -; 1.
DR InterPro; IPR005227; Cons_hypoth250.
DR InterPro; IPR006641; YqgFc.
DR Pfam; PF03652; UFP0081; 1.
DR SMART; SM00732; YqgFc; 1.
DR TIGRFAMs; TIGR00250; TIGR00250; 1.
KW Hydrolyase; Nuclease; DNA repair; DNA recombination; Complete proteome.
SQ SEQUENCE 156 AA; 17142 MW; 21F54D8648396141 CRC64;

Query Match 56.9%; Score 41; DB 1; Length 156;
Best Local Similarity 80.0%; Pred. No. 9.4;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGEQ 12
Db 18 AVVGLDPGEK 27

RESULT 4
MER_METTI
ID_MER_METTI STANDARD; PRT; 326 AA.
AC Q9UXP0;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Coenzyme F420-dependent N(5),N(10)-methylentetrahydromethanopterin
DE reductase (EC 1.5.99.11) (Methylene-H(4)MPT reductase).
GN MER OR FFD.
OS Methanobolus tindarius.
OC Archaea; Euryarchaeota; Methanomicrobia; Methanosarcinales;
OC Methanosarcinaceae; Methanobolus.
OX NCBI_TaxID=2221;
RN [1]
SEQUENCE FROM N.A.
RP STRAIN=DSM 2278;
RC MEDLINE=99132696; PubMed=9933933;
RX Westenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,
RA Gottschalk G., Blaut M.;
RT "The F420H2-dehydrogenase from Methanobolus tindarius: cloning of the
RT ffd operon and expression of the genes in Escherichia coli.";
RL FEMS Microbiol. Lett. 170:389-398(1999).
CC -!- FUNCTION: Catalyzes the reversible reduction of methylene-H(4)MPT
CC to methyl-H(4)MPT (By similarity).
CC -!- CATALYTIC ACTIVITY: N(5),N(10)-methylentetrahydromethanopterin +
CC reduced coenzyme F420 = 5-methyl-5,6,7,8-tetrahydromethanopterin +
CC coenzyme F420.
CC -!- PATHWAY: Methanogenesis from carbon dioxide; fifth step.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the mer family.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AJ011519; CAB56639.1; -.
DR PIR; T45226; T45226.
DR HAMAP; MF 01091; -; 1.
DR InterPro; IPR002103; Bac_luciferase.
DR Pfam; PF00296; bac_luciferase; 1.
KW Mechanogenesis; One-carbon metabolism; Oxidoreductase.
SQ SEQUENCE 326 AA; 34043 MW; 16F3AB97334A5D82 CRC64;

Query Match 56.9%; Score 41; DB 1; Length 326;
Best Local Similarity 70.0%; Pred. No. 19;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGEQ 12
Db 88 ALLGLGPGEQ 97

RESULT 5
COBI_MYCTU
ID_COBI_MYCTU STANDARD; PRT; 508 AA.
AC Q10677;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cobalamin biosynthesis protein COBIJ [Includes: Precorrin-2 C20-
DE methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2
DE methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.-)].
GN COBIJ OR COBI OR RV2066 OR MT2126 OR MTCY49.05 OR MB2092.
OS Mycobacterium tuberculosis, and
OS Mycobacterium bovis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773, 1765;
RN [1]
SEQUENCE FROM N.A.
RP SPECIES=M.tuberculosis; STRAIN=H37Rv;
RC MEDLINE=98295987; PubMed=9634230;
RX Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badoock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulten J.E., Taylor K., Whitehead S., Barrall B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
RN [2]
SEQUENCE FROM N.A.
RP SPECIES=M.tuberculosis; STRAIN=CDC 1551 / Oshkosh;
RC MEDLINE=22206494; PubMed=12218036;
RX Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;
RT "Whole-genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains.";
RL J. Bacteriol. 184:5479-5490(2002).
RN [3]
SEQUENCE FROM N.A.
RP SPECIES=M.bovis; STRAIN=AF2122/97;
RC MEDLINE=22709107; PubMed=12788972;
RX Garnier T., Eiglmeier K., Camus J.-C., Medina N., Mansoor H.,
RA Pryor M., Duthoy S., Grondin S., Lacroix C., Mounsepe C., Simon S.,
RA Harris B., Atkin R., Doggett J., Mayes R., Keating L., Wheeler P.R.,
RA Parkhill J., Barrall B.G., Cole S.T., Gordon S.V., Hewinson R.G.;
RT "The complete genome sequence of Mycobacterium bovis.";
RL Proc. Natl. Acad. Sci. U.S.A. 100:7877-7882(2003).
CC -!- FUNCTION: METHYLATES PRECORRIN-2 AT THE C-20 POSITION TO PRODUCE
CC PRECORRIN-3A (BY SIMILARITY).
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + precorrin-2 = S-
CC adenosyl-L-homocysteine + precorrin-3A.
CC -!- PATHWAY: Cobalamin biosynthesis.
CC -!- SIMILARITY: BELONGS TO A FAMILY THAT GROUPS SUMT, CYSG, CBIF/COBM
CC AND CBIL/COBI.
CC
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CC

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CC EMBL; Z73966; CAA98214.1; -
DR EMBL; AE007063; RAK46406.1; -
DR EMBL; BX248341; CAD96945.1; -
DR PIR; E70764; E70764.
DR TIGR; WP2126; -
DR Tuberculist; RV2066; -
DR InterPro; IPR006364; Cobi_CbiL.
DR InterPro; IPR006363; Cobi.
DR InterPro; IPR000878; Cor/por Metransf.
DR InterPro; IPR003043; Uropor Metransf.
DR Pfam; PR00590; TP_methylase_2.
DR TIGRPFAM; TIGR01457; cobi_cbiL; 1.
DR TIGRPFAM; TIGR01466; cobi_cbiH; 1.
DR PROSITE; PS00839; SUMT_1; 1.
DR PROSITE; PS00840; SUMT_2; 1.
KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW Methyltransferase; Multifunctional enzyme; Complete proteome.
FT DOMAIN 1 243 PRECORIN-2 C20-METHYLTRANSFERASE.
FT DOMAIN 244 508 PRECORIN-3 METHYLASE.
SQ SEQUENCE 508 AA; 53910 MW; 95AC066F02C24DC1 CRC64;

Query Match 56.9%; Score 41; DB 1; Length 508;
Best Local Similarity 58.3%; Pred. No. 30;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 AVVGLSPGQEY 14
Db 250 AVVGLGPGSDW 261
||||| ||: ||

RESULT 6
PLB4 SCHPO STANDARD; PRT; 673 AA.
AC Q9P327;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Putative lysophospholipase C977.09c precursor (EC 3.1.1.5)
DE (Phospholipase B).
GN SPAC977.09C OR SPAC1348.10C.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21848401; PubMed=11859360;
RA Wood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgouros J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule L., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volkart G., Aert R., Robben J., Grymonprez B.,
RA Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritzc C., Holzer E., Moestl D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Leirach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,

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RA Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe.";
RL Nature 415:871-880(2002)
CC -!- FUNCTION: Catalyzes the release of fatty acids from
CC lysophospholipids (by similarity).
CC -!- CATALYTIC ACTIVITY: 2-lysophosphatidylcholine + H(2)O =
CC glycerophosphocholine + a fatty acid anion.
CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
CC -!- SIMILARITY: Belongs to the lysophospholipase family.
CC
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CC
CC EMBL; AL358912; CAB94277.1; -
CC EMBL; AL137130; CAB69631.1; -
CC PIR; T50281; T50281.
CC GenBank SPORDB; SPAC977.09c; -
CC InterPro; IPR001179; FKSP_PPase.
CC InterPro; IPR002642; PLAC.
CC Pfam; PF01735; PLA2_B; 1.
CC SMART; SM00022; PLAC; 1.
KW Hypothetical protein; Lipid degradation; Hydrolase; Glycoprotein;
KW Signal.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 673 PUTATIVE LYSOPHOSPHOLIPASE C977.09C.
FT CARBOHYD 72 72 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 125 125 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 191 191 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 194 194 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 272 272 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 301 301 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 374 374 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 404 404 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 409 409 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 481 481 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 516 516 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 545 545 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 574 574 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 673 AA; 74595 MW; B39A773E76CD694B CRC64;

Query Match 55.6%; Score 40; DB 1; Length 673;
Best Local Similarity 61.5%; Pred. No. 58;
Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 2 PAVVGLSPGQEY 14
Db 83 PASEGLNEGEQSY 95
||||| ||: |||

RESULT 7
KV31 HUMAN STANDARD; PRT; 115 AA.
ID KV31 HUMAN
AC P04433;
DT 13-AUG-1987 (Rel. 05, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Ig kappa chain V-JII region VG precursor (fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=85087932; PubMed=6440122;
RA Pech M., Zachau H.G.;
RA "Immunoglobulin genes of different subgroups are interdigitated
RA within the VK locus.";
RA Nucleic Acids Res. 12:9229-9236(1984).

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CC -----
CC EMBL; X01668; -; NOT_ANNOTATED_CDS.
CC PIR; A01900; K3HUVG.
CC HSSP; P80362; 1WTL.
CC GO; GO:0005576; C:extracellular; NAS.
CC GO; GO:0003823; F:antigen binding; NAS.
CC GO; GO:0006955; P:immune response; NAS.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003596; Ig_v.
CC Pfam; PF00047; Ig; 1.
CC SMART; SM00406; IGV; 1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC Immunoglobulin V region; Signal.
KW SIGNAL 1 20 IG KAPPA CHAIN V-III REGION VG.
FT CHAIN 21 >115 FRAMEWORK-1.
FT DOMAIN 21 43 COMPLEMENTARITY-DETERMINING-1.
FT DOMAIN 44 54 FRAMEWORK-2.
FT DOMAIN 55 69 COMPLEMENTARITY-DETERMINING-2.
FT DOMAIN 70 76 FRAMEWORK-3.
FT DOMAIN 77 108 COMPLEMENTARITY-DETERMINING-3.
FT DOMAIN 109 115 BY SIMILARITY.
FT DISULFID 43 108
FT NON_TER 115
SQ SEQUENCE 115 AA; 12575 MW; 2DB47CDA3A17D555 CRC64;

Query Match 54.2%; Score 39; DB 1; Length 115;
Best Local Similarity 63.6%; Pred. No. 15;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12
DB 28 PAVVGLSPGEQ 38

RESULT A
KV5I_MOUSE STANDARD; PRT; 115 AA.
ID KV5I_MOUSE
AC P01642;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Ig kappa chain V-V region L7 precursor (Fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=81220975; PubMed=6264318;
RA Pech M., Hochti J., Schnell H., Zachau H.G.;
RT "Differences between germ-line and rearranged immunoglobulin V kappa
RL coding sequences suggest a localized mutation mechanism."
RL Nature 291:668-670(1981).
CC -1- MISCELLANEOUS: THERE APPEAR TO BE TWO POSSIBLE SPICE JUNCTIONS AT
CC THE 3' END OF THE INTRON. THE ALTERNATE WOULD CODE FOR A PROTEIN
CC LACKING RESIDUES 17-19.
CC PIR; A01925; KVM5L7.
CC PDB; 1J10; 18-FEB-03.
CC PDB; 1J1P; 18-FEB-03.
CC PDB; 1J1X; 18-FEB-03.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003596; Ig_v.
CC Pfam; PF00047; Ig; 1.
CC SMART; SM00406; IGV; 1.
CC PROSITE; PS50835; IG_LIKE; 1.

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KW Immunoglobulin V region; Signal; 3D-structure.
FT SIGNAL 1 20 IG KAPPA CHAIN V-V REGION L7.
FT CHAIN 21 >115 FRAMEWORK-1.
FT DOMAIN 21 43 COMPLEMENTARITY-DETERMINING-1.
FT DOMAIN 44 54 FRAMEWORK-2.
FT DOMAIN 55 69 COMPLEMENTARITY-DETERMINING-2.
FT DOMAIN 70 76 FRAMEWORK-3.
FT DOMAIN 77 108 COMPLEMENTARITY-DETERMINING-3.
FT DOMAIN 109 >115 BY SIMILARITY.
FT DISULFID 43 108
FT NON_TER 115
SQ SEQUENCE 115 AA; 12615 MW; C17BEC758C577E00 CRC64;

Query Match 54.2%; Score 39; DB 1; Length 115;
Best Local Similarity 54.5%; Pred. No. 15;
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12
DB 28 PAVVGLSPGEQ 38

RESULT 9
PVRU_HAEIN STANDARD; PRT; 278 AA.
ID PVRU_HAEIN
AC Q03432;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Formyltetrahydrofolate deformylase (EC 3.5.1.10) (Formyl-FH(4)
DE hydrolase).
GN PVRU OR H11588.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Haemophilus.
OX NCBI_TaxID=727;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=Rd / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kierkavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Utterback T.B., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RT "Whole-genome random sequencing and assembly of Haemophilus influenzae
RL Rd."
RL Science 269:496-512(1995).
RN [2]
RP SEQUENCE OF 64-278 FROM N.A.
RX STRAIN=RM 7004 / Serotype B;
RX MEDLINE=93328119; PubMed=8335255;
RA Maskell D.J.;
RT "Cloning and sequencing of the Haemophilus influenzae aroA gene."
RL Gene 129:155-156(1993).
CC -1- FUNCTION: PRODUCES FORMATE FROM FORMYL-TETRAHYDROFOLATE. PROVIDES
CC THE MAJOR SOURCE OF FORMATE FOR THE PURT-DEPENDENT SYNTHESIS OF
CC 5'-PHOSPHORIBOSYL-N-FORMYLGLYCINAMIDE (FGAR) DURING AEROBIC
CC GROWTH. HAS A ROLE IN REGULATING THE ONE-CARBON POOL
CC (BY SIMILARITY).
CC -1- CATALYTIC ACTIVITY: 10-formyltetrahydrofolate + H(2)O = formate +
CC tetrahydrofolate.
CC -1- ENZYME REGULATION: Activated by methionine, inhibited by glycine
CC (By similarity).
CC -1- PATHWAY: De novo purine biosynthesis.
CC -1- SUBUNIT: Homohexamer (By similarity).
CC -1- SIMILARITY: SOME, TO GAR TRANSFORMYLASE (PURN).
CC -----
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Orphan nuclear receptor HMR (Nerve growth factor induced protein I-B) (NGFI-B) (NUR77).

DR NRAA1 OR HMR OR NGFI-B.

OS Rattus norvegicus (Rat).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OX NCBI\_TaxID=10116;

RN SEQUENCE FROM N.A.

RP MEDLINE=90166506; PubMed=3272167;

RA Milbrandt J.;

RT "Nerve growth factor induces a gene homologous to the glucocorticoid receptor gene.";

RL Neuron 1:183-188(1986).

RN [2]

RP CHARACTERIZATION.

RX MEDLINE=93361012; PubMed=8995013;

RA Wilson T.E., Fahrner T.J., Milbrandt J.;

RT "The orphan receptors NGFI-B and steroidogenic factor 1 establish monomer binding as a third paradigm of nuclear receptor-DNA interaction.";

RL Mol. Cell. Biol. 13:5794-5804(1993).

RN [3]

RP DNA BINDING MOTIFS.

RX MEDLINE=92229411; PubMed=1314418;

RA Wilson T.E., Paulsen R.E., Padgett K.A., Milbrandt J.;

RT "Participation of non-zinc finger residues in DNA binding by two nuclear orphan receptors.";

RL Science 256:107-110(1992).

RN [4]

RP PHOSPHORYLATION.

RX MEDLINE=94043340; PubMed=8227042;

RA Hirata Y., Kiuchi K., Chen H.-C., Milbrandt J., Guroff G.;

RT "The phosphorylation and DNA binding of the DNA-binding domain of the orphan nuclear receptor NGFI-B.";

RL J. Biol. Chem. 268:24808-24812(1993).

RN [5]

RP X-RAY CRYSTALLOGRAPHY (2.7 ANGSTROMS) OF 264-351 IN COMPLEX WITH NBRE, AND METAL-BINDING.

RX MEDLINE=99260743; PubMed=10331876;

RA Meinke G., Sigler P.B.;

RT "DNA-binding mechanism of the monomeric orphan nuclear receptor NGFI-B.";

RL Nat. Struct. Biol. 6:471-477(1999).

CC -!- FUNCTION: Probable nuclear receptor. May act concomitantly with NUR1 in regulating the expression of delayed-early genes during liver regeneration. Binds the NGFI-B response element (NBRE) 5'-AAAAGGTCA-3'.

CC -!- SUBUNIT: Binds DNA as a monomer.

CC -!- SUBCELLULAR LOCATION: Nuclear.

CC -!- TISSUE SPECIFICITY: Expressed in lung, brain and superior cervical ganglia. High levels are seen in the adrenal tissue.

CC -!- INDUCTION: By nerve growth factor and during liver regeneration.

CC -!- PTM: Phosphorylation of Ser-350 results in decrease in NBRE binding while phosphorylation of Ser-340 has little effect on it.

CC -!- SIMILARITY: Belongs to the nuclear hormone receptor family. NR4 subfamily.

CC

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CC

CC EMBL; U17254; AAA56770.1; ALT\_INIT.

CC PDB; 1C1T; 26-JUN-00.

DR TRANSFAC; T00619; -

DR InterPro; IPR000536; Hormone\_rec\_lig.

DR InterPro; IPR001723; Strhmn\_receptor.

DR InterPro; IPR008946; Str\_ncl\_receptor.

DR InterPro; IPR001628; Znf\_C4steroid.

PFam; PF00104; hormone\_rec; 1.

DR PFam; PF00105; zF-C4; 1.

DR PRINTS; PRO0398; STRDHORMONER.

DR PRINTS; PRO0047; STROIDFINGER.

DR ProDom; PRO000035; Znf\_C4steroid; 1.

DR SMART; SM00430; HOLI; 1.

DR SMART; SM00399; Znf\_C4; 1.

DR PROSITE; PS00031; NUCLEAR RECEPTOR; 1.

KW Receptor; Transcription regulation; DNA-binding; Nuclear protein; Zinc-finger; Phosphorylation; 3D-structure.

FT DNAS\_BIND 266 331 NUCLEAR RECEPTOR-TYPE.

FT ZN\_FING 266 286 C4-TYPE.

FT ZN\_FING 302 326 C4-TYPE.

FT DOMAIN 408 458 LIGAND-BINDING (POTENTIAL).

FT DOMAIN 80 91 POLY-SER.

FT DOMAIN 182 186 POLY-PRO.

FT DOMAIN 582 585 POLY-PRO.

FT MOD\_RES 340 340 PHOSPHORYLATION (BY PKA).

FT MOD\_RES 350 350 PHOSPHORYLATION (BY PKA).

FT MUTAGEN 340 340 S->A: LOSS OF PHOSPHORYLATION.

FT MUTAGEN 350 350 S->A: LOSS OF PHOSPHORYLATION.

FT MUTAGEN 345 345 R->K: DECREASED NBRE BINDING.

FT MUTAGEN 348 348 L->V: ALMOST COMPLETE LOSS OF NBRE BINDING.

FT SEQUENCE 597 AA; 64281 MW; 9CFA987112337E53 CRC64;

SQ

Query Match 54.2%; Score 39; DB 1; Length 597;

Best Local Similarity 46.2%; Pred. No. 75;

Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPAVVGLSPGQEQE 13

Db 423 IPGFIELSPGDQD 435

RESULT 12

N160 MOUSE

ID N160 MOUSE STANDARD; PRT; 1402 AA.

AC Q9Z0W3; Q9CZD3;

DT 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Nuclear pore complex protein Nup160 (Nucleoporin Nup160) (160 kDa DE nucleoporin) (Gene trap locus 1-13) (GTL-13).

GN NUP160 OR GTL1-13 OR KIAA0197.

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

OX NCBI\_TaxID=10090;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=129/SVJ;

RA Van de Putte T., Cozijneen M., Dewulf N., Tylzanowski P., Lonnoy O., RA Huybroeck D.;

RT "Mus musculus mRNA for gtl-13 (gene trap locus-13), similar to human RT KIAA0197 gene (DB3781), complete cds.";

RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.

RN [2]

RP SEQUENCE OF 1151-1402 FROM N.A.

RC STRAIN=C57BL/6J; TISSUE=Embryo;

RX MEDLINE=21085660; PubMed=11217851;

RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y., RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S., RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I., RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R., RA Kadoya K., Matsuoka H.A., Ashburner M., Batalov S., Casavant T., RA Fleischmann W., Gaasterland T., Giesi C., King B., Kochiwa H., RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J., RA Schriml L.M., Stauble F., Suzuki R., Tomita M., Wagner L., Washio T., RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G., RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.P., RA Brownstein M.J., Sult C., Fletcher C., Fujita M., Gariboldi M., RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,

RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,  
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,  
 RA Sasaki H., Sato K., Schoenbach C., Seya I., Shibata Y., Storch K.-F.,  
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,  
 RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawai H., Kohtsuki S.,  
 RA Hayaishizaki Y.,  
 RA "Functional annotation of a full-length mouse cDNA collection.";  
 RL Nature 409:685-690(2001).  
 [3]  
 RP IDENTIFICATION, SUBUNIT, AND SUBCELLULAR LOCATION.  
 RX MEDLINE=21448620; PubMed=11564755;  
 RA Belagere N., Rabut G., Bai S.W., van Overbeek M., Beaudouin J.,  
 RA Daigle N., Zatepina O.V., Pasteau F., Labas V., Fromont-Racine M.,  
 RA Ellenberg J., Doye V.,  
 RA "An evolutionarily conserved NPC subcomplex, which redistributes in  
 part to kinetochores in mammalian cells.";  
 RL J. Cell Biol. 154:11147-11160(2001).  
 [4]  
 RP IDENTIFICATION, FUNCTION, SUBUNIT, AND SUBCELLULAR LOCATION.  
 RX MEDLINE=21541555; PubMed=11684705;  
 RA Vasu S., Shah S., Orjalo A., Park M., Fischer W.H., Forbes D.J.,  
 RA "Novel vertebrate nucleoporins Nup133 and Nup160 play a role in mRNA  
 export.";  
 RL J. Cell Biol. 155:339-354(2001).  
 CC -!- SUBUNIT: Involved in poly(A)+ RNA transport.  
 CC -!- SUBUNIT: Forms part of the Nup160 subcomplex in the nuclear pore  
 which is composed of Nup160, Nup133, Nup107 and Nup96. This  
 complex plays a role in RNA export and in tethering Nup98 and  
 Nup153 to the nucleus.  
 CC -!- SUBCELLULAR LOCATION: Nuclear pore complex.  
 CC -!- CAUTION: Ref.2 sequence differs from that shown due to a  
 frameshift in position 1157 and a stop codon in position 1396.  
 CC  
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 CC  
 CC EMBL; AF104415; AAD17922.2;  
 CC EMBL; AK012715; BAB28429.1; ALT\_FRAME.  
 CC GCD; MGI:1926227; Nup160.  
 CC GO; GO:0005643; C:nuclear pore; IDA.  
 CC GO; GO:0005487; F:nucleocytoplasmic transporter activity; IDA.  
 CC GO; GO:0006406; P:mRNA-nucleus export; IDA.  
 KW Nuclear protein; Transport.  
 FT CONFLICT 1156 1156 A -> T (IN REF. 2).  
 FT CONFLICT 1314 1314 E -> G (IN REF. 2).  
 FT CONFLICT 1368 1368 N -> D (IN REF. 2).  
 SQ SEQUENCE 1402 AA; 158230 MW; 3BF5DF057D28772 CRC64;  
 Query Match 54.2%; Score 39; DB 1; Length 1402;  
 Best Local Similarity 70.0%; Pred. No. 1.7e+02;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 LPAVVGSLSPG 10  
 Dd 170 IPSVFGSLSPG 179  
 RESULT 13  
 BUD3 YEAST  
 ID\_BUD3 YEAST STANDARD; PRT; 1636 AA.  
 AC P25556; P25557; P87007;  
 DT 01-MAY-1992 (Rel. 22, Created)  
 DT 01-NOV-1995 (Rel. 32, Last sequence update)  
 DT 10-OCT-2003 (Rel. 42, Last annotation update)  
 DE Bud site selection protein BUD3.  
 GN BUD3 OR YCL014W/YCL013W/YCL012W OR YCL14W/YCL13W/YCL12W.  
 OS Saccharomyces cerevisiae (Baker's yeast).  
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;

Saccharomycetales; Saccharomycetaceae; Saccharomycetes.  
 NCBI\_TaxID=4932;  
 [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=95247824; PubMed=7730410;  
 RA Chant J., Mischke M., Mitchell E., Herskowitz I., Pringle J.R.;  
 RA "Role of Bud3p in producing the axial budding pattern of yeast.";  
 RL J. Cell Biol. 129:767-778(1995).  
 [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=S288c;  
 RX MEDLINE=92244356; PubMed=1574125;  
 RA Oliver S.G.; van der Aart Q.J.M., Agostoni-Carbone M.L., Aigle M.,  
 RA Alberghina L., Alexandraki D., Antoine G., Anwar R., Ballesta J.P.G.,  
 RA Benit P., Berben G., Bergantino E., Biteau N., Bolle P.-A.,  
 RA Carlotin-Fukuhara M., Brown A.J.P., Brown R., Buhler J.-M.,  
 RA Carignani G., Chanet R., Contreras R., Crouzet M., Daignan-Fornier B.,  
 RA De Haan M., Defoor E., Delgado M.D., Demolder J., Doira C., Dubois E.,  
 RA Dujon B., Duesterhoeft A., Erdmann D., Esteban M., Fabre F.,  
 RA Fairhead C.A., Faye G., Feldmann H., Fiers W.,  
 RA Francinques-Gaillard M.-C., Franco L., Frontali L., Fukuhara H.,  
 RA Fuller L.J., Gent M.E., Gigot D., Gilliquet V., Glandsdorff N.,  
 RA Goffeau A., Grenson M., Grisanti P., Grivell L.A., Haesemaun M.,  
 RA Hatat D., Hegemann J.H., Herbert C.J., Hilger F., Hohmann S.,  
 RA Hollenberg C.P., Huse K., Iborra F., Indge K.J., Isono K., Jackman P.,  
 RA Jacq C., Jacquet M., James C.M., Jauniaux J.-C., Jia Y., Jimenez A.,  
 RA Kleinhaus U., Kreisel P., Lafranchi G., Lewis C., van der Linden C.G.,  
 RA Lucchini G., Kutzekirchen K., Maat C., Manhaupt M., Manzano M.B.,  
 RA Lucchini G., Kutzekirchen K., Maat C., Manhaupt M., Manzano M.B.,  
 RA Martegani E., Mathieu A., Maurer C.T.C., McConnell D., McKee R.A.,  
 RA Messenguy F., Mewes H.-W., Molemans F., Montague M.A., Navas L.,  
 RA Newton C.S., Olson M.V., Pallier C., Panzeri L., Pearson B.M.,  
 RA Pereira J., Philippsen P., Pierard A., Planta R.J., Plevani P.,  
 RA Poetsch B., Pohl F.M., Purnelle B., Ramezani Rad M., Rasmussen S.W.,  
 RA Raynal A., Remacha M., Richterich P., Roberts A.B., Rodriguez F.,  
 RA Sanz E., Schaaff-Gerstenschlaeger I., Scherens B., Schweitzer B.,  
 RA Shu Y., Skala J., Slonimski P.P., Sor F., Soustelle C., Thierry A.,  
 RA Spiegelberg R., Statera L.I., Steensma H.F., Steiner S., Vetter I.,  
 RA Thiegs G., Triano L.N., Urrestazu L.A., Valle G., Vetter I.,  
 RA van Vliet-Reedijk J.C., Volckaert G., Vreken P., Warrington J.R.,  
 RA von Wettstein D., Wicksteed B.L., Wilson C., Wurst H., Xu G.,  
 RA Zimmermann F.K., Sgourou J.G.,  
 RT "The complete DNA sequence of yeast chromosome III.";  
 RL Nature 357:38-46(1992).  
 [3]  
 RP REVISIONS.  
 RA Gromadka R.;  
 RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.  
 [4]  
 RP REVISIONS.  
 RA Valles G., Volckaerts G.;  
 RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
 CC -!- FUNCTION: Co-assembles with BUD4 at bud sites. BUD4 and BUD3 may  
 cooperate to recognize a spatial landmark (the neck filaments)  
 during mitosis and they subsequently become a landmark for  
 establishing the axial budding pattern in G1.  
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 CC  
 CC EMBL; U17580; AAA86315.1;  
 CC EMBL; X59720; CAA42346.2;  
 DR PIR; S74285; S74285.  
 DR PIR; S74286; S74286.  
 DR Germonline; 138855;  
 DR SGD; S0000520; BUD3.  
 DR InterPro; IPR000219; RhoGEF.  
 DR SMART; SM00325; RhoGEF; 1.  
 KW Cell cycle.

```

SQ SEQUENCE 1636 AA; 184717 MW; 9E4E45BA5C3A3P69 CRC64;
Query Match 54.2%; Score 39; DB 1; Length 1636;
Best Local Similarity 58.3%; Pred. No. 2e+02;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 PAVVGLSPGQEQ 13
||: |||: ||
Db 112 PAIENLSPSQEQ 123

RESULT 14
COBL_MYCTU STANDARD; PRT; 390 AA.
ID COBL_MYCTU
AC Q10671;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Precorrin-6Y C5,15-methyltransferase [decarboxylating] (EC 2.1.1.132)
DE (Precorrin-6 methyltransferase) (Precorrin-6Y methylase).
DE COBL OR RV2072C OR MT2132 OR MTCY49.11C.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN SEQUENCE FROM N.A.
RP STRAIN=H37RV;
RC MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holtroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RX MEDLINE=22206494; PubMed=12218036;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,
RA Delcher A., Uterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;
RA "Whole-genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains.";
RL J. Bacteriol. 184:5479-5490 (2002).
CC -!- FUNCTION: CATALYZES THE METHYLATION OF BOTH C-5 AND C-15 IN
CC PRECORRIN-6Y TO FORM PRECORRIN-8X.
CC -!- CATALYTIC ACTIVITY: 2 S-adenosyl-L-methionine + precorrin-6Y = 2
CC S-adenosyl-L-homocysteine + precorrin-8X + CO(2).
CC -!- PATHWAY: Cobalamin biosynthesis.
CC -!- SIMILARITY: TO S-TYRIMURUM CBIE; ALSO, LOW, TO OTHER
CC METHYLASES INVOLVED IN COBALAMIN BIOSYNTHESIS.
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CC
CC EMBL; 273966; CAA38225.1; -.
CC DR EMBL; AE007063; AAK46412.1; -.
CC PIR; C70765; C70765.
CC DR TIGR; MT2132; -.
CC TubercuList; RV2072c; -.

DR InterPro; IPR006365; COBL.
DR InterPro; IPR000878; Cor/pox Metransf.
DR InterPro; IPR000051; SAM_bind.
DR Pfam; PF00590; TP_methylase; 1.
DR TIGRFAMS; TIGR01468; cobL cbiet; 1.
KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW Methyltransferase; Complete proteome.
FT CONFLICT 205 L -> P (IN REF. 2).
FT CONFLICT 327 327 D -> H (IN REF. 2).
SQ SEQUENCE 390 AA; 41854 MW; FB42EFF7562F21F3 CRC64;

Query Match 52.8%; Score 38; DB 1; Length 390;
Best Local Similarity 88.9%; Pred. No. 73;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 LPAVVGGLSP 9
|||: |||: |||
Db 55 LPAVQGLSP 63

RESULT 15
RSI_LEULA STANDARD; PRT; 429 AA.
ID RSI_LEULA
AC P50889; P71450;
DT 01-OCT-1996 (Rel. 34, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE 40S ribosomal protein S1.
GN RPS1.
OS Leuconostoc lactis.
OC Bacteria; Firmicutes; Lactobacillales; Leuconostoc.
OX NCBI_TaxID=1246;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97186703; PubMed=9034319;
RA Yamit-Hezi A., Levy Z., Neuman S., Nudel U.;
RA "A Leuconostoc lactis protein with homology to ribosomal protein S1
RT shares common epitopes and common DNA binding properties with a
RT mammalian DNA binding nuclear factor.";
RL Gene 185:99-103 (1997).
RN [2]
RP SEQUENCE OF 24-429 FROM N.A.
RX MEDLINE=95237615; PubMed=7721096;
RA Eklund E.A., Lee S.W., Skalniak D.G.;
RT "Cloning of a cDNA encoding a human DNA-binding protein similar to
RL ribosomal protein S1.";
RN Gene 155:231-235 (1995).
RP [3]
RP SEQUENCE OF 78-429 FROM N.A.
RX MEDLINE=96164600; PubMed=8568274;
RA Tsuzaka K., Leu A.K., Frank M.B., Movafagh B.F., Koscec M.,
RA Winkler T.H., Kalden J.R., Reichlin M.;
RT "Lupus autoantibodies to double-stranded DNA cross-react with
RL ribosomal protein S1.";
RN J. Immunol. 156:1668-1675 (1996).
CC -!- FUNCTION: EXHIBITS PREFERENTIAL BINDING TO SINGLE-STRANDED AND
CC DOUBLE-STRANDED DNA AND A LOW BINDING AFFINITY FOR RNA.
CC -!- SIMILARITY: Belongs to the S1P family of ribosomal proteins.
CC -!- SIMILARITY: Contains 4 S1 motif domains.
CC -!- CAUTION: WAS ORIGINALLY (REF.2 AND REF.3) THOUGHT TO ORIGINATE
CC FROM HUMAN BUT IS MOST PROBABLY THE RESULT OF A CDNA LIBRARY
CC CONTAMINATION BY L.LACTIS.
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CC
CC EMBL; U24086; AAB08978.1; -.
CC DR EMBL; U05589; AAA77669.1; -.

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DR EMBL; U27517; AAA97575.1; -.
DR HSP; P05055; LSRO.
DR InterPro; IPR008994; Nucleic acid OB.
DR InterPro; IPR00110; Ribosomal_S1.
DR InterPro; IPR003029; S1.
DR Pfam; PF00575; S1; 4.
DR PRINTS; PR00681; RIBOSOMALS1.
DR SMART; SM00316; S1; 4.
DR PROSITE; PS0126; S1; 4.
KW Ribosomal protein; Repeat; RNA-binding.
FT DOMAIN 55 128 S1 MOTIF 1.
FT DOMAIN 144 211 S1 MOTIF 2.
FT DOMAIN 231 299 S1 MOTIF 3.
FT DOMAIN 316 385 S1 MOTIF 4.
FT CONFLICT 24 24 S -> G (IN REF. 2).
FT CONFLICT 122 122 A -> S (IN REF. 3).
FT CONFLICT 217 217 L -> R (IN REF. 2 AND 3).
SQ SEQUENCE 429 AA; 46386 MW; 92AC82605F39DDFC CRC64;

Query Match      52.8%; Score 38; DB 1; Length 429;
Best Local Similarity 80.0%; Pred.No. 80;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 3 AVVGLSPGEQ 12
Db 71 AVVGLSTGEE 80

```

Search completed: May 7, 2004, 12:34:33  
 Job time : 7.04 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:26:40 ; Search time 26.18 Seconds  
(without alignments)  
168.726 Million cell updates/sec

Title: US-09-786-214A-12

Perfect score: 72

Sequence: 1 LPAVVGSLSPGEQY 14

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:\*

- 1: sp\_archaea:\*
- 2: sp\_bacteria:\*
- 3: sp\_fungi:\*
- 4: sp\_human:\*
- 5: sp\_invertebrate:\*
- 6: sp\_mammal:\*
- 7: sp\_mhc:\*
- 8: sp\_organalle:\*
- 9: sp\_phage:\*
- 10: sp\_plant:\*
- 11: sp\_rodent:\*
- 12: sp\_virus:\*
- 13: sp\_vertebrate:\*
- 14: sp\_unclassified:\*
- 15: sp\_virus:\*
- 16: sp\_bacteriap:\*
- 17: sp\_archaeap:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	49	68.1	41	11	Q8K408
2	44	61.1	381	16	Q9RL5
3	43.5	60.4	614	16	Q8Y280
4	43	59.7	821	17	Q9HPR8
5	42	58.3	1541	6	Q8HXL3
6	41	56.9	228	17	Q9HKE3
7	41	56.9	326	1	Q9UXP0
8	41	56.9	579	4	Q8N158
9	41	56.9	877	2	Q7X128
10	41	56.9	11096	2	Q9L4W3
11	40	55.6	154	17	Q8ZYJ3
12	40	55.6	227	17	Q97WD1
13	40	55.6	242	16	Q7WCX1
14	40	55.6	261	16	Q8NQA2
15	40	55.6	298	16	Q7W5D6
16	40	55.6	298	16	Q7VTI2

17	40	55.6	353	17	Q8ZU64
18	40	55.6	355	10	Q947A7
19	40	55.6	357	4	Q8NEX1
20	40	55.6	358	4	Q9NWD0
21	40	55.6	386	4	Q9NWM3
22	40	55.6	428	16	Q92AG9
23	40	55.6	428	16	Q8Y661
24	40	55.6	485	10	Q7XNE4
25	40	55.6	540	16	Q9RR71
26	40	55.6	661	10	Q8S8J8
27	40	55.6	665	16	Q8XR57
28	40	55.6	683	2	Q34302
29	40	55.6	753	16	Q89T31
30	39	54.2	148	16	Q8F4Y0
31	39	54.2	197	17	Q8FWI9
32	39	54.2	209	17	Q8TTT1
33	39	54.2	209	17	Q8TH81
34	39	54.2	277	16	Q9KQK6
35	39	54.2	277	16	Q87RD3
36	39	54.2	284	16	Q7VFI5
37	39	54.2	319	16	Q8ZKQ5
38	39	54.2	319	16	Q8Z2X4
39	39	54.2	407	2	Q9LCW0
40	39	54.2	589	16	Q7UPH0
41	39	54.2	626	16	Q7VWC3
42	39	54.2	627	16	Q7WH05
43	39	54.2	627	16	Q7W9Q3
44	39	54.2	1400	11	Q7TME1
45	38.5	53.5	94	16	Q88F51

#### ALIGNMENTS

##### RESULT 1

Q8K408 ID Q8K408 PRELIMINARY; PRT; 41 AA.  
AC Q8K408;  
DT 01-OCT-2002 (TREMBLrel. 22, Created)  
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)  
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)  
DE Truncated macrophage colony stimulating factor.  
GN CSF1.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
OX NCBI\_taxID=10116;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=LEW.tl;  
RX MEDLINE=22069908; PubMed=12074592;  
RA Dobbins D.E., Sood R., Hashiramoto A., Hansen C.T., Wilder R.L., Remmers E.F.;  
RA "Mutation of macrophage colony stimulating factor (CSF1) causes osteopetrosis in the tl rat.";  
RT Biochem. Biophys. Res. Commun. 294:1114-1120(2002).  
DR EMBL; AF514357; AAM54137.1; -.  
SQ SEQUENCE 41 AA; 4178 MW; 1D342C19BD18AA41 CRC64;

Query Match 68.1%; Score 49; DB 11; Length 41;

Best Local Similarity 76.9%; Pred. No. 0.46; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGSLSPGEQ 13

DB 21 LPAAGLSPEQ 33

##### RESULT 2

Q9RL5 ID Q9RL5 PRELIMINARY; PRT; 381 AA.

AC Q9RL5; Q7W5D6

DT 01-MAY-2000 (TREMBLrel. 13, Created)



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DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DE 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Cytochrome P450.
GN DR2473.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
RX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heideberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Utterback T., Zalewski C.,
RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RT "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1.";
RL Science 286:1571-1577(1999).
CC -!- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.
DR EMBL; AF002076; AAF12016.1; -.
DR PIR; F75270; F75270.
DR TIGR; DR2473; -.
DR GO; GO:0004497; F:monooxygenase activity; IEA.
DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR001128; Cytochrome_P450.
DR Pfam; PF00067; P450; 1.
DR PRINTS; PR00385; P450.
DR PROSITE; PS00086; CYTOCHROME_P450; 1.
KW Heme; Monooxygenase; Oxidoreductase; Complete proteome.
SQ SEQUENCE 381 AA; 41940 MW; F191EA69F1797B53 CRC64;

Query Match 51.1%; Score 44; DB 16; Length 381;
Best Local Similarity 100.0%; Pred. No. 37;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGSLSP 9
DB 52 LPAVVGSLSP 60

ID Q8Y280 PRELIMINARY; PRT; 614 AA.
AC Q8Y280;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Probable ATP-binding transport ABC transporter protein.
GN RSC0456 OR RS04444.
OS Ralstonia solanacearum (Pseudomonas solanacearum).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Burkholderiaceae; Ralstonia.
OX NCBI_TaxID=305;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=GM11000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
RA Arlat M., Billault A., Brottier P., Camus J.C., Catolico L.,
RA Chandler M., Choïene N., Claude-Renard C., Cunnac S., Denange N.,
RA Gaspin C., Lavie M., Moisan A., Robert C., Saurin W., Schiex T.,
RA Sigüier P., Thebault P., Whalen M., Wincker P., Levy M.,
RA Weissenbach J., Boucher C.A.;
RT "Genome sequence of the plant pathogen Ralstonia solanacearum.";
RL Nature 415:497-502(2002).
DR EMBL; AL646059; CAD13984.1; -.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0004009; F:ATP-binding cassette (ABC) transporter acti. . .; IEA.

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DR GO; GO:0000166; F:nucleotide binding; IEA.
DR GO; GO:0006810; P:transport; IEA.
DR InterPro; IPR003593; AAA_ATPase.
DR InterPro; IPR003439; ABC_transporter.
DR InterPro; IPR001623; DnaJ_N.
DR Pfam; PF00005; ABC_tran; 1.
DR ProDom; PD0000006; ABC_transporter; 1.
DR SMART; SM00382; AAA_1.
DR PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
DR PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
DR PROSITE; PS00636; DnaJ_1; 1.
KW Complete proteome.
SQ SEQUENCE 614 AA; 69240 MW; E293355B85872142 CRC64;

Query Match 60.4%; Score 43.5; DB 16; Length 614;
Best Local Similarity 45.8%; Pred. No. 76;
Matches 11; Conservative 2; Mismatches 0; Indels 11; Gaps 1;

QY 1 LPAVVG-----LSPGEEQ 13
DB 501 LPAVGLLDEVSNWSLRLSPGEEQ 524

ID Q9HPR8 PRELIMINARY; PRT; 821 AA.
AC Q9HPR8;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE DNA helicase.
GN HEL OR VNG1501G.
OS Halobacterium sp. (strain NRC-1 / ATCC 700922 / JCM 11081).
OC Archaea; Euryarchaeota; Halobacteria; Halobacteriales;
OC Halobacteriaceae; Halobacterium.
OX NCBI_TaxID=64091;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=20504483; PubMed=11016950;
RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,
RA Shukla H.D., Lasky S.R., Balliga N.S., Thorsson V., Sbrogna J.,
RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
RA Leithauser B., Keller K., Cruz R., Danson M.J., Hough D.W.,
RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
RA Isenbarger T.A., Peck R.F., Pohlschroder M., Spudich J.L., Jung K.-H.,
RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
RA Ebhardt H., Lowe T.M., Liang P., Riley M., Hood L., DasSarma S.;
RT "Genome sequence of Halobacterium species NRC-1.";
RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).
DR EMBL; AE005064; AAG19799.1; -.
DR PIR; C84304; C84304.
DR GO; GO:0004386; F:helicase activity; IEA.
DR GO; GO:0000166; F:nucleotide binding; IEA.
DR GO; GO:0007242; P:intracellular signalling cascade; IEA.
DR InterPro; IPR003593; AAA_ATPase.
DR InterPro; IPR001478; PDZ.
DR SMART; SM00382; AAA; 1.
DR SMART; SM00228; PDZ; 1.
KW Helicase; Complete proteome.
SQ SEQUENCE 821 AA; 89848 MW; C454C76B984A5702 CRC64;

Query Match 59.7%; Score 43; DB 17; Length 821;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGEEQ 12
DB 326 AVVGLSPAEEQ 335

ID Q8HXL3 PRELIMINARY; PRT; 1541 AA.

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AC Q9HXL3; 2003 (TrEMBLrel. 23, Created)  
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Hypothetical protein.  
 OS Sus scrofa (pig).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
 OX NCBI\_TaxID=9823;  
 RN [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=21845885; PubMed=11856879;  
 RA Drogemuller C., Kuiper H., Voss-Nemitz R., Brenig B., Distl O.,  
 RA Leeb T.;  
 RT "Molecular characterization and chromosome assignment of the porcine  
 RT gene COX7A1 coding for the muscle specific cytochrome c oxidase  
 RT subunit VIIa-M";  
 RT subunit VIIa-M";  
 RL CytoGenet. Cell Genet. 94:190-193 (2001).  
 RN [2]  
 RN  
 RP SEQUENCE FROM N.A.  
 RA Leeb T.;  
 RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ410870; CAD56046.1; -;  
 DR InterPro; IPR001680; WD40.  
 DR Pfam; PF00400; WD40; 12.  
 DR PROSITE; PS00082; WD\_REPEATS\_2; 1.  
 DR PROSITE; PS02994; WD\_REPEATS\_REGION; 3.  
 KW Hypothetical protein.  
 SQ SEQUENCE 1541 AA; 168898 MW; 81B8882854FAF4F1E CRC64;  
 Query Match 58.3%; Score 42; DB 6; Length 1541;  
 Best Local Similarity 58.3%; Pred. No. 3.7e+02;  
 Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 1 LPAVVLSPGGEQ 12  
 : : : |||||  
 Db 786 MFSEISLSPGGEQ 797

RESULT 6  
 Q9HKE3 PRELIMINARY; PRT; 228 AA.  
 AC Q9HKE3;  
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)  
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Precorrin-2 methyltransferase related protein.  
 GN TA0658.  
 OS Thermoplasma acidophilum.  
 OC Archaea; Euryarchaeota; Thermoplasmatia; Thermoplasmatales;  
 OC Thermoplasmataceae; Thermoplasma.  
 OX NCBI\_TaxID=2303;  
 RN [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RX STRAIN=DSM 1728;  
 RX MEDLINE=20479972; PubMed=11029001;  
 RA Ruepp A., Graml W., Santos-Martinez M.-L., Koretke K.K., Volker C.,  
 RA Mewes H.-W., Frishman D., Stocker S., Lupas A.N., Baumeister W.;  
 RT "The genome sequence of the thermoacidophilic scavenger Thermoplasma  
 RT acidophilum.";  
 RL Nature 407:508-513 (2000).  
 DR EMBL; AL445065; CAC11796.1; -;  
 DR GO; GO:0008757; F:S-adenosylmethionine-dependent methyltransf. . .; IEA.  
 DR GO; GO:0016740; F:transferase activity; IEA.  
 DR GO; GO:0008152; P:metabolism; IEA.  
 DR GO; GO:0006779; P:porphyrin biosynthesis; IEA.  
 DR InterPro; IPR006364; Cobi CbiL.  
 DR InterPro; IPR000878; Cor/For\_Mettransf.  
 DR InterPro; IPR003043; Uropor\_Mettransf.  
 DR Pfam; PF00590; TP\_methylase; 1.  
 DR TIGRFAMs; TIGR01467; cobi cbiL; 1.  
 DR PROSITE; PS00839; SUMT 1; 1.  
 DR Transferase; Methyltransferase; Complete proteome.

SQ SEQUENCE 228 AA; 25084 MW; 11ABD8B68192A67C CRC64;  
 Query Match 56.9%; Score 41; DB 17; Length 228;  
 Best Local Similarity 63.6%; Pred. No. 70;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 4 VVGLSPGGEQ 14  
 : : : |||||  
 Db 5 VVGLSPGGEQ 15

RESULT 7  
 Q9UXP0 PRELIMINARY; PRT; 326 AA.  
 AC Q9UXP0;  
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)  
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE F420-dependent N5,N10-methylene-tetrahydromethanopterin reductase,  
 DE putative.  
 GN FPDA.  
 OS Methanobolus tindarius.  
 OC Archaea; Euryarchaeota; Euryarchaeota orders incertae sedis;  
 OC Methanosarcinales; Methanosarcinaceae; Methanobolus.  
 OX NCBI\_TaxID=2221;  
 RN [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=DSM 2278;  
 RA Westenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,  
 RA Gottschalk G., Blaut M.;  
 RT "The F420H2-dehydrogenase from Methanobolus tindarius: Cloning of the  
 RT ffd operon and expression of the genes in Escherichia coli.";  
 RL Submitted (SEP-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ011519; CAB56639.1; -;  
 DR PIR; T45226; T45226.  
 DR InterPro; IPR002103; Bac\_luciferase.  
 DR Pfam; PF00296; bac\_luciferase; 1.  
 SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;  
 Query Match 56.9%; Score 41; DB 1; Length 326;  
 Best Local Similarity 70.0%; Pred. No. 1e+02;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 3 AVVGLSPGGEQ 12  
 : : : |||||  
 Db 88 AILGLSPGGEQ 97

RESULT 8  
 Q8N158 PRELIMINARY; PRT; 579 AA.  
 ID Q8N158;  
 AC Q8N158;  
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)  
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Similar to cerebroglycan (Hypothetical protein FLJ38962).  
 GN DKFZP547M109.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 OX NCBI\_TaxID=9606;  
 RN [1]  
 RN  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Brain;  
 RA Strausberg R.;  
 RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RN  
 RP SEQUENCE FROM N.A.  
 RA Ninomiya K., Wagatsuma M., Kanda K., Kondo H., Yokoi T., Kodaira H.,  
 RA Furuya T., Takahashi M., Kikkawa E., Omura Y., Abe K., Kamiyama K.,  
 RA Katsuta N., Sato K., Tanikawa M., Yamazaki M., Sugiyama T., Irie R.,  
 RA Otsuki T., Sato H., Wakamatsu A., Ishii S., Yamamoto J., Isono Y.,  
 RA Kawai-Hio Y., Saito K., Nishikawa T., Kimura K., Yamashita H.,

Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
Streptomycineae; Streptomycetaceae; Streptomycetes.  
NCBI\_TaxID=1971;  
[1]  
RN  
RP SEQUENCE FROM N.A.  
RC STRAIN=ATCC 11455;  
RX MEDLINE=20334850; PubMed=10873841;  
RA Brautaet T., Sekurova O.N., Sletta H., Ellingsen T.E., Strom A.R.,  
RA Valla S., Zotchev S.B.;  
RT "Biosynthesis of the polyene antifungal antibiotic nystatin in  
RT Streptomyces noursei ATCC 11455: analysis of the gene cluster and  
RT deduction of the biosynthetic pathway.";  
RL Chem. Biol. 7:395-403 (2000).  
DR EMBL; AF2633912; RAF1776.1; -;  
DR HSSP; P25715; IMLA.  
DR GO; GO:0004024; F:alcohol dehydrogenase activity, zinc-dependent; IEA.  
DR GO; GO:0016740; F:transferase activity; IEA.  
DR GO; GO:0008270; F:zinc ion binding; IEA.  
DR GO; GO:0004314; F:[acyl-carrier protein] S-malonyltransferase. . ; IEA.  
DR GO; GO:0006633; P:fatty acid biosynthesis; IEA.  
DR GO; GO:0008152; P:metabolism; IEA.  
DR InterPro; IPR001227; AC trans.  
DR InterPro; IPR002865; Adh\_zn\_family.  
DR InterPro; IPR004410; Fabb.  
DR InterPro; IPR000794; Ketoacyl synth.  
DR InterPro; IPR006162; Ppantne S.  
DR InterPro; IPR006163; Pp.bind.  
DR Pfam; PF00698; Acyl transf; 6.  
DR Pfam; PF00107; ADH\_Zinc\_N; 1.  
DR Pfam; PF00109; ketoacyl-synt; 6.  
DR Pfam; PF02801; ketoacyl-synt\_C; 6.  
DR Pfam; PF00550; pp-binding; 6.  
DR TIGRfams; TIGR00128; fabb; 6.  
DR PROSITE; PS00075; ACP DOMAIN; 6.  
DR PROSITE; PS00606; B\_KETOACYL SYNTHASE; 6.  
DR PROSITE; PS00012; PHOSPHOPANTETHEINE; 5.  
DR Phosphopantetheine; Transferase.  
SQ SEQUENCE 11096 AA; 1150415 MW; 776CAFAFCAE551DD CRC64;  
KW  
Query Match 56.9%; Score 41; DB 2; Length 11096;  
Best Local Similarity 63.6%; Pred.No. 4.7e+03;  
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps  
QY 2 PAVVGLSPGEQ 12  
DB 4998 PEVTGLPGDQ 5008  
RESULT 11  
Q8ZVJ3 PRELMINARY; PRT; 154 AA.  
AC Q8ZVJ3;  
AD 01-WAR-2002 (TrEMBLrel. 20, Created)  
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)  
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
DE Hypothetical protein PAE0746.  
GN PAE0746.  
OS Pyrobaculum aerophilum.  
OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;  
OX Thermoproteaceae; Pyrobaculum.  
NCBI\_TaxID=13773;  
[1]  
RN SEQUENCE FROM N.A.  
RC STRAIN=IM2 / ATCC 51768 / DSM 7523;  
RX MEDLINE=21664397; PubMed=11792869;  
RA Fitz-Gibbon S.T., Lubner H., Kim U.-J., Stetter K.O., Simon M.I.,  
RA Miller J.H.;  
RT "Genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum  
RT aerophilum";  
RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989(2002) .  
RE EMBL; AB009783; AAL63000.1; -;  
RW Hypothetical protein; Complete proteome.  
SQ SEQUENCE 154 AA; 17403 MW; 1C3D8BCB40324766 CRC64;  
KW

Query Match 55.6%; Score 40; DB 17; Length 154;  
 Best Local Similarity 66.7%; Pred. No. 68;  
 Matches 10; Conservative 1; Mismatches 2; Indels 2; Gaps 1;

QY 1 LPVVVGLS--PGEQE 13  
 |||: |||: |||:  
 Db 32 LPDDVGISYTPGEQE 46  
 |||: |||: |||:

RESULT 12  
 Q97WD1 PRELIMINARY; PRT; 227 AA.  
 AC Q97WD1;  
 DT 01-OCT-2001 (TRENBLrel. 18, Created)  
 DT 01-OCT-2001 (TRENBLrel. 18, Last sequence update)  
 DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)  
 DE Cobalamin biosynthesis precorrin-6B methylase, putative (cblE).  
 GN CBIIE OR SSO2296.  
 OS Sulfolobus solfataricus.  
 OC Archaea; Crenarchaeota; Thermoprotei; Sulfolobales; Sulfolobaceae;  
 OC Sulfolobus.  
 OX NCBI\_TaxID=2287;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 35092 / DSM 1617 / P2;  
 RX MEDLINE=21332296; Pubmed=11427726;  
 RA She Q., Singh R.K., Confalonieri F., Zivanovic Y., Allard G.,  
 RA Awayez M.J., Chan-Weiher C.C.-Y., Clausen I.G., Curtis B.A.,  
 RA De Moors A., Erasuo G., Fletcher C., Gordon P.M.K.,  
 RA Heikamp-de Jong I., Jeffries A.C., Kozera C.J., Medina N., Peng X.,  
 RA Thi-Ngoc H.P., Redder P., Schenk M.E., Theriault C., Tolstrup N.,  
 RA Charlebois R.L., Doolittle W.F., Duguet M., Gaasterland T.,  
 RA Garrett R.A., Ragan M.A., Sensen C.W., Van der Oost J.;  
 RA "The complete genome of the crenarchaeon Sulfolobus solfataricus P2.";  
 RT Proc. Natl. Acad. Sci. U.S.A. 98:7835-7840(2001).  
 DR EMBL; AB006833; AAK42457.1; -;  
 DR PIR; B90400.  
 DR GO; GO:0008168; P:methyltransferase activity; IEA.  
 DR GO; GO:0008152; P:metabolism; IEA.  
 DR InterPro; IPR000878; Cor/por\_Mettransf.  
 DR Pfam; PF00590; TP\_methylase; 1.  
 KW Methyltransferase; Complete proteome.  
 SQ SEQUENCE 227 AA; 25548 MW; 0267F09491F2DCED CRC64;

Query Match 55.6%; Score 40; DB 17; Length 227;  
 Best Local Similarity 54.5%; Pred. No. 1e+02;  
 Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 4 VVGLSPGGEQEY 14  
 :||: ||: ||:  
 Db 10 IVGVGPDPEY 20  
 :||: ||: ||:

RESULT 13  
 Q7WCX1 PRELIMINARY; PRT; 242 AA.  
 AC Q7WCX1;  
 DT 01-OCT-2003 (TRENBLrel. 25, Created)  
 DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)  
 DE Bacteriophage-related DNA polymerase.  
 GN B3808.  
 OS Bordetella bronchiseptica (Alcaligenes bronchisepticus).  
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;  
 OC Alcaligenaceae; Bordetella.  
 OX NCBI\_TaxID=518;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=RB50 / ATCC BAA-588;  
 RX MEDLINE=22827954; Pubmed=12910271;  
 RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,

RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,  
 RA Cerdano-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,  
 RA Achtman M., Atkin R., Baker S., Basham D., Bason N., Cherevach I.,  
 RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,  
 RA Feltwell T., Goble A., Hanlin N., Hauser H., Holroyd S., Jagels K.,  
 RA Leather S., Moule S., Norberczak H., O'Neill S., Ormond D., Price C.,  
 RA Rabinowitsch E., Rutter S., Sanders M., Saunders D., Seeger K.,  
 RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,  
 RA Unwin I., Whitehead S., Barrell B.G., Maskell D.J.;  
 RT "Comparative analysis of the genome sequences of Bordetella pertussis,  
 RT Bordetella parapertussis and Bordetella bronchiseptica.";  
 RT Nat. Genet. 35:32-40(2003).  
 DR EMBL; BX640448; CAE35782.1; -;  
 KW Complete proteome.  
 SQ SEQUENCE 242 AA; 25627 MW; EECAD9B319823284 CRC64;

Query Match 55.6%; Score 40; DB 16; Length 242;  
 Best Local Similarity 80.0%; Pred. No. 1.1e+02;  
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGGEQE 13  
 |||: |||: |||:  
 Db 96 VVGEAPGGEQE 105  
 |||: |||: |||:

RESULT 14  
 Q8NQA2 PRELIMINARY; PRT; 261 AA.  
 AC Q8NQA2;  
 DT 01-OCT-2002 (TRENBLrel. 22, Created)  
 DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)  
 DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)  
 DE Hypothetical protein Cgl1536.  
 GN CG11536.  
 OS Corynebacterium glutamicum (Brevibacterium flavum).  
 OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
 OC Corynebacterineae; Corynebacteriaceae; Corynebacterium.  
 OX NCBI\_TaxID=1718;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;  
 RA Nakagawa S.;  
 RT "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";  
 RL Submitted (MAY-2002) to the EMBL/GenBank/DBSJ databases.  
 DR EMBL; AP005278; BAB98929.1; -;  
 KW Hypothetical protein; Complete proteome.  
 SQ SEQUENCE 261 AA; 27957 MW; 4D8D51D4DCA3A210 CRC64;

Query Match 55.6%; Score 40; DB 16; Length 261;  
 Best Local Similarity 53.8%; Pred. No. 1.2e+02;  
 Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPVVVGLSPGGEQE 13  
 |||: |||: |||:  
 Db 137 LPATVSPGEAD 149  
 |||: |||: |||:

RESULT 15  
 Q7W5D6 PRELIMINARY; PRT; 298 AA.  
 AC Q7W5D6;  
 DT 01-OCT-2003 (TRENBLrel. 25, Created)  
 DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)  
 DE Bacteriophage-related DNA polymerase.  
 GN BPP3357.  
 OS Bordetella parapertussis.  
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;  
 OC Alcaligenaceae; Bordetella.  
 OX NCBI\_TaxID=519;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=12822 / ATCC BAA-587;

```

RX MEDLINE=22827954; PubMed=12910271;
RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,
RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,
RA Cerdano-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,
RA Achtman M., Atkin R., Baker S., Basham D., Bason N., Cherevach I.,
RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,
RA Feltwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
RA Leather S., Moule S., Norberczak H., O'Neil S., Ormond D., Price C.,
RA Rabinowitsch E., Rutter S., Sanders M., Saunders D., Seeger K.,
RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,
RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;
RT "Comparative analysis of the genome sequences of Bordetella pertussis,
RT Bordetella parapertussis and Bordetella bronchiseptica.";
RL Nat. Genet. 35:32-40(2003);
DR EMBL; BX640433; CAE38642.1; -.
KW Complete proteome.
SQ SEQUENCE 298 AA; 31202 MW; A858C8199E2AC8B7 CRC64;

Query Match 55.6%; Score 40; DB 16; Length 298;
Best Local Similarity 80.0%; Pred. No. 1.4e+02;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGEQE 13
Db 152 VVGEAPGEQE 161
||| :|||
||| :|||

```

Search completed: May 7, 2004, 12:37:54  
Job time : 28.3467 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 39.06 Seconds  
(without alignments)  
101.272 Million cell updates/sec

Title: US-09-786-214A-12

Perfect score: 72

Sequence: 1 LPAVGLSPGEQY 14

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A Genesep29Jan04.\*

1: Genesep1980s.\*

2: Genesep1990s.\*

3: Genesep2000s.\*

4: Genesep2001s.\*

5: Genesep2002s.\*

6: Genesep2003as.\*

7: Genesep2003bs.\*

8: Genesep2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	72	100.0	14	3	AY84266 Peptide d
2	72	100.0	20	3	AY84265 Truncated
3	72	100.0	25	3	AY84264 Peptide o
4	68	94.4	13	3	AY84267 Peptide d
5	65	90.3	13	3	AY84268 Peptide d
6	65	90.3	15	3	AY84269 Peptide d
7	45	62.5	234	4	AAB36208 Human imm
8	42	58.3	470	6	AAB34724 Streptomy
9	42	58.3	475	6	AAB34732 Streptomy
10	42	58.3	475	6	AAB34729 Streptomy
11	41	56.9	106	5	ABJ10397 Mutant an
12	41	56.9	106	5	ABJ10395 Mutant an
13	41	56.9	306	4	ABG19552 Novel hum
14	41	56.9	530	7	ADD49105 Human NOV
15	41	56.9	549	7	ADD49091 Human NOV
16	41	56.9	579	5	ABG70277 Human Gly
17	41	56.9	579	5	ABG97356 Human GCD
18	41	56.9	579	6	ABR39111 Human GPC
19	41	56.9	579	7	ADD49087 Human NOV
20	41	56.9	579	7	ADD49107 Human NOV
21	41	56.9	579	7	ADD49089 Human NOV
22	41	56.9	592	7	ADD49099 Human NOV
23	41	56.9	11096	4	AAE10129 Streptomy
24	40	55.6	261	4	AAG91441 C glutami
25	40	55.6	287	6	ABU17217 Protein e

26	40	55.6	303	6	ADA36829	Acinetoba
27	40	55.6	306	4	ABG24698	Novel hum
28	40	55.6	358	4	AAB92530	Human pro
29	40	55.6	359	5	ABB97563	Novel hum
30	40	55.6	428	5	ABB47705	Listeria
31	40	55.6	668	3	AAB57055	Human pro
32	39	54.2	18	2	AA41875	Rheumatoi
33	39	54.2	18	4	AAU25388	Schizophr
34	39	54.2	18	4	AAU15732	Schizophr
35	39	54.2	18	5	ABG78871	Multiple
36	39	54.2	23	5	ABP62618	Human imm
37	39	54.2	74	6	ABJ18694	Antibody
38	39	54.2	88	3	AA556655	Partial p
39	39	54.2	93	5	AAU80983	Human ant
40	39	54.2	94	7	ADD69248	Human lig
41	39	54.2	95	6	ABO27154	Human ger
42	39	54.2	95	6	ABO27153	Human ger
43	39	54.2	96	6	ABO27155	Human ger
44	39	54.2	96	6	ABO27150	Human ger
45	39	54.2	100	5	AAE23987	Human MOG

## ALIGNMENTS

### RESULT 1

AY84266

ID AAY84266 standard; peptide; 14 AA.

XX

AC AAY84266;

XX

DT 12-JUL-2000 (first entry)

XX

DE Peptide derived from macrophage colony stimulating gene alternative ORF.

XX

KW tumour rejection antigen; macrophage colony stimulating gene;

KW macrophage-colony stimulating factor; antigen presenting cell;

KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX

OS Synthetic.

OS Homo sapiens.

XX

FN WO200013699-A1.

XX

PD 16-MAR-2000.

XX

PF 03-SEP-1999; 99WO-US020344.

XX

PR 04-SEP-1998; 98US-0099077P.

XX

PA (LUDW-) LUDWIG INST CANCER RES.

XX

PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX

DR WPI; 2000-256859/22.

XX

PT Isolated polypeptide used to treat subjects having a disorder characterized by expression of alternative open reading frame macrophage-colony stimulating factor comprises 25 amino acid residue sequence.

XX

Claim 2; Page 39; 7app; English.

XX

CC The present sequence represents a peptide which is derived from a tumour rejection antigen precursor encoded by an alternative open reading frame (ORF) of human macrophage colony stimulating gene. Peptides derived from the alternative ORF of macrophage-colony stimulating factor, when presented by an antigen presenting cell having a human leukocyte antigen (HLA) class I molecule, effectively induce the activation and proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF of macrophage-colony stimulating factor are useful for enriching selectively a population of T lymphocytes with CD8+ T lymphocytes. They are also useful for diagnosing a disorder characterized by expression of the polypeptide, and for identifying



KW macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 OS Synthetic.  
 OS Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 PD 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 PI WPI; 2000-256859/22.  
 DR Isolated polypeptide used to treat subjects having a disorder  
 XX characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 PT Example 2; Page 40; 74pp; English.  
 PS The present sequence represents a peptide which is derived from a tumour  
 XX rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 94.4%; Score 68; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.00026; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 PAVVGLSPGEQY 14  
 DB 1 PAVVGLSPGEQY 13  
 |||||  
 RESULT 5  
 AAY84268  
 ID AAY84268 standard; peptide; 13 AA.  
 AC AAY84268;  
 XX 12-JUL-2000 (first entry)  
 DT Peptide derived from macrophage colony stimulating gene alternative ORF.  
 DE tumour rejection antigen; macrophage colony stimulating gene;  
 DE macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 KW Synthetic.  
 OS Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 PD 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 PI WPI; 2000-256859/22.  
 DR Isolated polypeptide used to treat subjects having a disorder  
 XX characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 PT Example 2; Page 40; 74pp; English.  
 PS The present sequence represents a peptide which is derived from a tumour  
 XX rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 94.4%; Score 68; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.00026; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 PAVVGLSPGEQY 14  
 DB 1 PAVVGLSPGEQY 13  
 |||||  
 RESULT 5  
 AAY84268  
 ID AAY84268 standard; peptide; 13 AA.  
 AC AAY84268;  
 XX 12-JUL-2000 (first entry)  
 DT Peptide derived from macrophage colony stimulating gene alternative ORF.  
 DE tumour rejection antigen; macrophage colony stimulating gene;  
 DE macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 KW Synthetic.  
 OS Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 PD 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 PI WPI; 2000-256859/22.  
 DR Isolated polypeptide used to treat subjects having a disorder  
 XX characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 PT Example 2; Page 40; 74pp; English.  
 PS The present sequence represents a peptide which is derived from a tumour  
 XX rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 90.3%; Score 65; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0008; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LPVVGLSPGEQE 13  
 DB 1 LPVVGLSPGEQE 13  
 |||||  
 RESULT 6  
 AAY84269  
 ID AAY84269 standard; peptide; 15 AA.  
 AC AAY84269;  
 XX 12-JUL-2000 (first entry)  
 DT Peptide derived from macrophage colony stimulating gene alternative ORF.  
 DE tumour rejection antigen; macrophage colony stimulating gene;  
 DE macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 KW Synthetic.  
 OS Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 PD 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 PI WPI; 2000-256859/22.  
 DR Isolated polypeptide used to treat subjects having a disorder  
 XX characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 PT Example 2; Page 40; 74pp; English.  
 PS The present sequence represents a peptide which is derived from a tumour  
 XX rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 90.3%; Score 65; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0008; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LPVVGLSPGEQE 13  
 DB 1 LPVVGLSPGEQE 13  
 |||||

PR 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 XX WPI; 2000-256859/22.  
 XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 XX Example 2; Page 40; 74pp; English.  
 XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 90.3%; Score 65; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0008; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LPVVGLSPGEQE 13  
 DB 1 LPVVGLSPGEQE 13  
 |||||  
 RESULT 6  
 AAY84269  
 ID AAY84269 standard; peptide; 15 AA.  
 AC AAY84269;  
 XX 12-JUL-2000 (first entry)  
 DT Peptide derived from macrophage colony stimulating gene alternative ORF.  
 DE tumour rejection antigen; macrophage colony stimulating gene;  
 DE macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
 KW Synthetic.  
 OS Homo sapiens.  
 XX WO200013699-A1.  
 XX 16-MAR-2000.  
 PD 03-SEP-1999; 99WO-US020344.  
 XX 04-SEP-1998; 98US-0099077P.  
 XX (LUDW-) LUDWIG INST CANCER RES.  
 PA Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
 PI WPI; 2000-256859/22.  
 DR Isolated polypeptide used to treat subjects having a disorder  
 XX characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.  
 PT Example 2; Page 40; 74pp; English.  
 PS The present sequence represents a peptide which is derived from a tumour  
 XX rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX Sequence 13 AA;  
 SQ Query Match 90.3%; Score 65; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0008; Indels 0; Gaps 0;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LPVVGLSPGEQE 13  
 DB 1 LPVVGLSPGEQE 13  
 |||||



PS Example 2; Page 40; 74pp; English.

CC The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 15 AA;

Query Match 90.3%; Score 65; DB 3; Length 15;  
 Best Local Similarity 100.0%; Pred. No. 0.00094;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LPAVVGSLSPGEQ 13  
 |||||  
 Db 3 LPAVVGSLSPGEQ 15

RESULT 7

AAE36208  
 ID AAB36208 standard; protein; 234 AA.

AC AAB36208;

DT 15-FEB-2001 (first entry)

XX Human immune system associated protein HISAP-6.

DE Human; immune system associated protein; HISAP-6; immune disorder;  
 KW infection; autoimmune disease; cancer.

OS Homo sapiens.

XX US6135941-A.

XX 24-OCT-2000.

XX 27-MAR-1998; 98US-00049672.

XX 27-MAR-1998; 98US-00049672.

XX (INCY-) INCYTE PHARM INC.

PI Tang YT, Yue H, Lal P, Corley NC, Guegler KJ, Baughn MR;

PI Hillman JL, Au-Young J;

XX WPI; 2001-030926/04.

DR N-PSDB; AAC66524.

XX New human immune system associated proteins (HISAP) and polynucleotides  
 FT encoding the HISAP, useful for diagnosing, treating or preventing immune  
 FT or cell proliferative disorders or infections.

PS Claim 1; Col 59-60; 54pp; English.

XX The present invention provides the coding and protein sequences for a  
 CC number of human immune system associated proteins (HISAPs). These can be  
 CC used in the diagnosis and treatment of various autoimmune disorders,  
 CC infections and cell proliferation diseases. The diseases include AIDS,  
 CC adult respiratory distress syndrome, anaemia, asthma, atherosclerosis,  
 CC Crohn's disease, irritable bowel syndrome, multiple sclerosis, myasthenia  
 CC gravis, osteoarthritis, rheumatoid arthritis, scleroderma, systemic lupus  
 CC erythematosus, arteriosclerosis, cirrhosis and cancer

XX Sequence 234 AA;

Query Match 62.5%; Score 45; DB 4; Length 234;  
 Best Local Similarity 72.7%; Pred. No. 36;  
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 PAVVGLSPGEQ 12  
 ||:|||||  
 Db 28 PAIVSLSPGER 38

RESULT 8

AAE34724

ID AAE34724 standard; protein; 470 AA.

XX AAE34724;

DT 14-MAY-2003 (first entry)

DE Streptomyces rimosus ema3 protein.

XX P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
 KW emamectin; insecticide; ema3 protein.

OS Streptomyces rimosus.

XX WC200292801-A2.

XX 21-NOV-2002.

XX 15-MAY-2002; 2002WO-EP005363.

XX 16-MAY-2001; 2001US-0291149P.

XX (SYGN) SYNGENTA PARTICIPATIONS AG.

XX Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;

PI Buckel TG;

XX WPI; 2003-140280/13.

DR N-PSDB; AADS3019.

XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.

XX Claim 17; Page 107-108; 157pp; English.

XX The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus ema3 protein

XX Sequence 470 AA;

Query Match 58.3%; Score 42; DB 6; Length 470;  
 Best Local Similarity 72.7%; Pred. No. 2.4e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 1 LPAVVGSLSPGE 11  
 ||:|||||  
 Db 44 LPSYVGLHPGE 54

RESULT 9

AAE34732

ID AAE34732 standard; protein; 475 AA.

XX AAE34732;

XX

DT 14-MAY-2003 (first entry)  
 XX Streptomyces rimosus ema1 protein.  
 DE P450 monooxygenase; avermectin; ferroxidoxin; ferroxidoxin reductase; enzyme;  
 KW emamectin; insecticide; ema1 protein.  
 XX Streptomyces rimosus.  
 OS Streptomyces rimosus.  
 XX WO200292801-A2.  
 FN 21-NOV-2002.  
 XX 15-MAY-2002; 2002WO-EP005363.  
 PF 16-MAY-2001; 2001US-0291149P.  
 PR (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 XX Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; AAD53027.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 121-122; 157pp; English.  
 PS The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferroxidoxins and ferroxidoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus ema1 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 58.3%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. No. 2.4e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 ||: ||| |||  
 Db 49 LPSYVGLHPGE 59  
 RESULT 10  
 ID AAE34729 standard; protein; 475 AA.  
 XX AAE34729;  
 AC AAE34729;  
 XX 14-MAY-2003 (first entry)  
 DT Streptomyces albofaciens ema8 protein.  
 DE P450 monooxygenase; avermectin; ferroxidoxin; ferroxidoxin reductase; enzyme;  
 KW emamectin; insecticide; ema8 protein.  
 XX Streptomyces albofaciens.  
 OS Streptomyces albofaciens.  
 XX WO200292801-A2.  
 FN 21-NOV-2002.  
 PD 15-MAY-2002; 2002WO-EP005363.  
 PF 16-MAY-2001; 2001US-0291149P.  
 PR

XX (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 XX Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; RAD53024.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 116-117; 157pp; English.  
 PS The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferroxidoxins and ferroxidoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces albofaciens ema8 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 58.3%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. No. 2.4e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 ||: ||| |||  
 Db 49 LPSYVGLHPGE 59  
 RESULT 11  
 ID ABJ10397 standard; protein; 106 AA.  
 XX ABJ10397;  
 AC ABJ10397;  
 XX 28-NOV-2002 (first entry)  
 DT Mutant anti-mesothelin Fv (ST6) variable light chain.  
 DE Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.  
 XX Unidentified.  
 OS Synthetic.  
 XX WO200240545-A2.  
 FN 23-MAY-2002.  
 PD 16-NOV-2001; 2001WO-US043602.  
 XX 17-NOV-2000; 2000US-0249805P.  
 PR (USSH ) US DEPT HEALTH & HUMAN SERVICES.  
 XX Pastan IH, Onda M, Nagata S, Tsutsui Y, Vincent JJ, Kreitman RJ;  
 PI Vasmatazis G, Lee B;  
 XX WPI; 2002-500208/53.  
 DR N-PSDB; AHT08085.  
 XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.  
 XX Claim 12; Fig 2; 82pp; English.

CC The invention comprises the amino acid and coding sequences of  
 CC recombinant immunotoxin proteins. The immunotoxin proteins of the  
 CC invention contain an antibody (or an antigen-binding fragment) with a  
 CC substitution of a negatively charged amino acid for an uncharged or  
 CC positively charged amino acid. The immunotoxins of the invention have  
 CC reduced liver toxicity. The immunotoxins of the invention are useful for  
 CC killing a malignant cell (e.g. a cancer cell). The present amino acid  
 CC sequence represents a recombinant immunotoxin of the invention  
 CC  
 XX Sequence 106 AA;

Query Match 56.9%; Score 41; DB 5; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 67;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 2 PAVVGLSPGEQ 12  
 Db 8 PAIMSASPGEQ 18  
 ||:|

RESULT 12  
 ABU10395  
 ID ABU10395 standard; protein; 106 AA.  
 XX  
 AC ABU10395;

XX 28-NOV-2002 (first entry)

DE Mutant anti-Tac Fv (M16) variable light chain.

XX Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.

XX Unidentified.  
 OS Synthetic.

PN WO200240545-A2.

XX 23-MAY-2002.

XX 16-NOV-2001; 2001WO-US043602.

XX 17-NOV-2000; 2000US-0249805P.

XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.

XX Pastan JH, Onda M, Nagata S, Tsutsumi Y, Vincent JJ, Kreitman RJ;  
 PI Vasmatazis G, Lee B;

XX WPI; 2002-500208/53.  
 DR N-PSDB; APT08083.

XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.

XX Claim 8; Fig 2; 82pp; English.

XX The invention comprises the amino acid and coding sequences of  
 CC recombinant immunotoxin proteins. The immunotoxin proteins of the  
 CC invention contain an antibody (or an antigen-binding fragment) with a  
 CC substitution of a negatively charged amino acid for an uncharged or  
 CC positively charged amino acid. The immunotoxins of the invention have  
 CC reduced liver toxicity. The immunotoxins of the invention are useful for  
 CC killing a malignant cell (e.g. a cancer cell). The present amino acid  
 CC sequence represents a recombinant immunotoxin of the invention  
 CC

XX Sequence 106 AA;

Query Match 56.9%; Score 41; DB 5; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 67;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 2 PAVVGLSPGEQ 12  
 Db 8 PAIMSASPGEQ 18  
 ||:|

RESULT 13  
 ABG19552  
 ID ABG19552 standard; protein; 306 AA.  
 XX  
 AC ABG19552;

XX 13-FEB-2002 (first entry)

DE Novel human diagnostic protein #19543.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;

XX food supplement; medical imaging; diagnostic; genetic disorder.

XX Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US008631.

XX 31-MAR-2000; 2000US-00540217.

XX 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX N-PSDB; AAS83739.

XX New isolated polynucleotide and encoded polypeptides, useful in

XX diagnostics, forensics, gene mapping, identification of mutations  
 XX responsible for genetic disorders or other traits and to assess  
 XX biodiversity.

XX Claim 20; SEQ ID NO 49911; 103pp; English.  
 XX The invention relates to isolated polynucleotide (I) and polypeptide (II)  
 CC sequences. (I) is useful as hybridisation probes, polymerase chain  
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,  
 CC and in recombinant production of (II). The polynucleotides are also used  
 CC in diagnostics as expressed sequence tags for identifying expressed  
 CC genes. (I) is useful in gene therapy techniques to restore normal  
 CC activity of (II) or to treat disease states involving (II). (II) is  
 CC useful for generating antibodies against it, detecting or quantitating a  
 CC polypeptide in tissue, as molecular weight markers and as a food  
 CC supplement. (II) and its binding partners are useful in medical imaging  
 CC of sites expressing (II). (I) and (II) are useful for treating disorders  
 CC involving aberrant protein expression or biological activity. The  
 CC polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity  
 CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. ABG0010-ABG30377 represent novel human diagnostic  
 CC amino acid sequences of the invention. Note: The sequence data for this  
 CC patent did not appear in the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 306 AA;

Query Match 56.9%; Score 41; DB 4; Length 306;  
 Best Local Similarity 57.1%; Pred. No. 2.2e+02;  
 Matches 8; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

Qy 1 LPVVGLSPGEQ 14  
 ||:|

CC epilepsy, immune disorders (osteoarthritis), hematopoietic disorders,  
 CC inflammatory skin disorders, asthma and various dyslipidemias. The coding  
 CC sequences and proteins may also be used as targets for the identification  
 CC of small molecules that modulate or inhibit e.g. neurogenesis, cell  
 CC differentiation, cell proliferation, haematopoiesis, wound healing and  
 CC angiogenesis, in gene therapy, in generation of antibodies that bind  
 CC immunospecifically to NOV substances for use in therapeutic or diagnostic  
 CC methods.  
 CC XX

SQ Sequence 530 AA;  
 Query Match 56.9%; Score 41; DB 7; Length 530;  
 Best Local Similarity 75.0%; Pred. No. 4e+02;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPVVVGLSPGQ 12  
 |||||  
 Db 416 LPVVVGGSPAEQ 427

RESULT 15  
 ADD49091

ID ADD49091 standard; protein; 549 AA.

XX AC ADD49091;

XX 15-JAN-2004 (first entry)

XX Human NOV15d SEQ ID 64.

XX Antidiabetic; anorectic; cardiast; hypotensive; antiarteriosclerotic;  
 XX viricide; antibacterial; fungicide; protozoacide; nootropic;  
 KW neuroprotective; anti-parkinsonian; anticonvulsant; osteopathic;  
 KW antiallergic; anti-inflammatory; dermatological; antiasthmatic;  
 KW antileptic; gene therapy; NOV protein; metabolic disorder; diabetes;  
 KW obesity; viral infection; bacterial infection; fungal infection;  
 KW helminthic infection; protozoal infection; anorexia; cancer;  
 KW cardiovascular disease; hypertension; atherosclerosis;  
 KW neurodegenerative disorder; Alzheimer's disease; Parkinson's disease;  
 KW epilepsy; immune disorder; osteoarthritis; haematopoietic disorder;  
 KW inflammatory skin disorder; asthma; dyslipidemia; human.

OS Homo sapiens.

XX WO2003060149-A2.

XX 24-JUL-2003.

XX 06-JAN-2003; 2003WO-US000252.

XX 04-JAN-2002; 2002US-0345222P.

XX 14-JAN-2002; 2002US-0348693P.

XX 16-JAN-2002; 2002US-0349182P.

XX 17-JAN-2002; 2002US-0349733P.

XX 18-JAN-2002; 2002US-0350263P.

XX 24-JAN-2002; 2002US-0351977P.

XX 28-MAY-2002; 2002US-0383758P.

XX 05-JUN-2002; 2002US-0385969P.

XX 11-JUN-2002; 2002US-0387834P.

XX 17-JUL-2002; 2002US-0396407P.

XX 30-SEP-2002; 2002US-0415115P.

XX 03-JAN-2003; 2003US-00336603.

XX (CURA-) CURAGEN CORP.

XX Grosse WM, Alsobrook JP, Anderson DW, Burgess CE, Edinger SR;  
 XX Ellerman K, Furtak K, Gangolli EA, Gerlach VL, Gilbert JA;  
 XX Gunther E, Gorman L, Guo X, Ji W, Li L, Miller CE, Padigar M;  
 XX Patturajan M, Rastelli L, Macdougall JR, Mishra VS, Smithson G;  
 XX Spytek KA, Stone DU, Shenoy SG, Taupier RJ, Vernet CAM, Zhong M;  
 XX Malyankar UM, Millet I, Kekuda R;  
 XX WPI; 2003-587288/55.

Db 137 LPQAPGLSPGAQSW 150

RESULT 14

ADD49105

ID ADD49105 standard; protein; 530 AA.

XX AC ADD49105;

XX 15-JAN-2004 (first entry)

XX Human NOV15k SEQ ID 78.

XX Antidiabetic; anorectic; cardiast; hypotensive; antiarteriosclerotic;  
 KW viricide; antibacterial; fungicide; protozoacide; nootropic;  
 KW neuroprotective; anti-parkinsonian; anticonvulsant; osteopathic;  
 KW antiallergic; anti-inflammatory; dermatological; antiasthmatic;  
 KW antileptic; gene therapy; NOV protein; metabolic disorder; diabetes;  
 KW obesity; viral infection; bacterial infection; fungal infection;  
 KW helminthic infection; protozoal infection; anorexia; cancer;  
 KW cardiovascular disease; hypertension; atherosclerosis;  
 KW neurodegenerative disorder; Alzheimer's disease; Parkinson's disease;  
 KW epilepsy; immune disorder; osteoarthritis; haematopoietic disorder;  
 KW inflammatory skin disorder; asthma; dyslipidemia; human.

XX Homo sapiens.

XX WO2003060149-A2.

XX 24-JUL-2003.

XX 06-JAN-2003; 2003WO-US000252.

XX 04-JAN-2002; 2002US-0345222P.

XX 14-JAN-2002; 2002US-0348693P.

XX 16-JAN-2002; 2002US-0349182P.

XX 17-JAN-2002; 2002US-0349733P.

XX 18-JAN-2002; 2002US-0350263P.

XX 24-JAN-2002; 2002US-0351977P.

XX 28-MAY-2002; 2002US-0383758P.

XX 05-JUN-2002; 2002US-0385969P.

XX 11-JUN-2002; 2002US-0387834P.

XX 17-JUL-2002; 2002US-0396407P.

XX 30-SEP-2002; 2002US-0415115P.

XX 03-JAN-2003; 2003US-00336603.

XX (CURA-) CURAGEN CORP.

XX Grosse WM, Alsobrook JP, Anderson DW, Burgess CE, Edinger SR;  
 XX Ellerman K, Furtak K, Gangolli EA, Gerlach VL, Gilbert JA;  
 XX Gunther E, Gorman L, Guo X, Ji W, Li L, Miller CE, Padigar M;  
 XX Patturajan M, Rastelli L, Macdougall JR, Mishra VS, Smithson G;  
 XX Spytek KA, Stone DU, Shenoy SG, Taupier RJ, Vernet CAM, Zhong M;  
 XX Malyankar UM, Millet I, Kekuda R;  
 XX WPI; 2003-587288/55.  
 XX N-ESDB; ADD49104.

XX New isolated NOVX polypeptides and polynucleotides, useful for

XX preventing, diagnosing or treating NOVX-associated disorders, e.g.

XX osteoarthritis, obesity, atherosclerosis, cancer, Parkinson's disease,

XX asthma, or infections.

XX Claim 1; Page 181-182; 31pp; English.

XX The present invention relates to novel NOV proteins and their coding

XX sequences (ADD49028-ADD49131). The proteins and coding sequences are

XX useful in the manufacture of a medicament for treating a syndrome

XX associated with a human disease, preferably a NOV-associated disorder

XX such as metabolic disorders, diabetes, obesity, infectious diseases

XX (viral, bacterial, fungal, helminthic, and protozoal), anorexia, cancer,

XX cardiovascular diseases (hypertension, atherosclerosis),

XX neurodegenerative disorders, Alzheimer's disease, Parkinson's disease,

DR N-PSDB; ADD49090.  
XX New isolated NOVX polypeptides and polynucleotides, useful for  
PT preventing, diagnosing or treating NOVX-associated disorders, e.g.  
PT osteoarthritis, obesity, atherosclerosis, cancer, Parkinson's disease,  
PT asthma, or infections.  
XX  
PS Claim 1; Page 177; 311pp; English.  
XX  
XX The present invention relates to novel NOV proteins and their coding  
CC sequences (ADD49028-ADD49131). The proteins and coding sequences are  
CC useful in the manufacture of a medicament for treating a syndrome  
CC associated with a human disease, preferably a NOV-associated disorder  
CC such as metabolic disorders, diabetes, obesity, infectious diseases  
CC (viral, bacterial, fungal, helminthic, and protozoal), anorexia, cancer,  
CC cardiovascular diseases (hypertension, atherosclerosis),  
CC neurodegenerative disorders (Alzheimer's disease, Parkinson's disease,  
CC epilepsy, immune disorders (osteoarthritis), hematopoietic disorders,  
CC inflammatory skin disorders, asthma and various dyslipidemias. The coding  
CC sequences and proteins may also be used as targets for the identification  
CC of small molecules that modulate or inhibit e.g. neurogenesis, cell  
CC differentiation, cell proliferation, hematopoiesis, wound healing and  
CC angiogenesis, in gene therapy, in generation of antibodies that bind  
CC immunospecifically to NOV substances for use in therapeutic or diagnostic  
CC methods.  
XX  
SQ Sequence 549 AA;  
  
Query Match 56.9%; Score 41; DB 7; Length 549;  
Best Local Similarity 75.0%; Pred. No. 4.1e+02;  
Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
  
QY 1 LPAVVGLSPGEQ 12  
Db 419 LPPVVGSPAEQ 430  
  
Search completed: May 7, 2004, 12:33:45  
Job time : 41.06 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:26:40 ; Search time 24.31 Seconds  
(without alignments)  
168.726 Million cell updates/sec

Title: US-09-786-214A-13  
Perfect score: 68  
Sequence: 1 PAVVGLSPGQEY 13

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

- 1: sp.archaea.\*
- 2: sp.bacteria.\*
- 3: sp.fungi.\*
- 4: sp.human.\*
- 5: sp.invertebrate.\*
- 6: sp.mammal.\*
- 7: sp.mhc.\*
- 8: sp.organelle.\*
- 9: sp.phage.\*
- 10: sp.plant.\*
- 11: sp.rodent.\*
- 12: sp.virus.\*
- 13: sp.vertebrate.\*
- 14: sp.unclassified.\*
- 15: sp.rvirus.\*
- 16: sp.bacteriap.\*
- 17: sp.archaeap.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	45	66.2	41	11 Q8K408	Q8K408 rattus norv
2	43	63.2	821	17 Q9HPR8	Q9hpr8 halobacteri
3	41	60.3	228	17 Q9HKE3	Q9hke3 thermoplas
4	41	60.3	326	1 Q9UXP0	Q9uxp0 methanolobu
5	41	60.3	877	2 Q7X128	Q7xi28 xanthomonas
6	41	60.3	11096	2 Q9L4W3	Q9l4w3 streptomyce
7	40	58.8	227	17 Q97WD1	Q97wd1 sulfolobus
8	40	58.8	242	16 Q7WCX1	Q7wxc1 bordetella
9	40	58.8	238	16 Q7W5D6	Q7w5d6 bordetella
10	40	58.8	298	16 Q7VTI2	Q7vti2 bordetella
11	40	58.8	355	10 Q947A7	Q947a7 nitellopsis
12	40	58.8	357	4 Q8NHX1	Q8nhx1 homo sapien
13	40	58.8	358	4 Q9NWD0	Q9nwd0 homo sapien
14	40	58.8	381	16 Q9RRL5	Q9rrl5 deinococcus
15	40	58.8	386	4 Q9NMW3	Q9nmw3 homo sapien
16	40	58.8	540	16 Q9RR71	Q9rr71 deinococcus

17	40	58.8	661	10 Q8S8J8	Q8s8j8 arabidopsis
18	40	58.8	685	16 Q8XR57	Q8xr57 ralstonia s
19	40	58.8	1541	6 Q8HXL3	Q8hxl3 sus scrofa
20	39.5	58.1	614	16 Q8Y280	Q8y280 ralstonia s
21	39	57.4	148	16 Q8FY0	Q8fy0 leptospira
22	39	57.4	197	17 Q8PW19	Q8pw19 methanosarc
23	39	57.4	209	17 Q8TTL1	Q8ttl1 methanosarc
24	39	57.4	209	17 Q8TH81	Q8th81 methanosarc
25	39	57.4	407	2 Q9LCW0	Q9lcw0 streptomyce
26	39	57.4	428	16 Q92AG9	Q92ag9 listeria in
27	39	57.4	428	16 Q8Y661	Q8y661 listeria mo
28	39	57.4	589	16 Q7UPH0	Q7uph0 rhodospirell
29	39	57.4	626	16 Q7VWC3	Q7vwc3 bordetella
30	39	57.4	626	16 Q7WH05	Q7wh05 bordetella
31	39	57.4	627	16 Q7W9Q3	Q7w9q3 bordetella
32	39	57.4	627	16 Q88F51	Q88f51 pseudomonas
33	38.5	56.6	94	16 Q88F51	Q81061 mus musculu
34	38	55.9	136	11 Q61061	Q9V2S2 pyrococcus
35	38	55.9	152	17 Q8TZK1	Q8tzk1 pyrococcus
36	38	55.9	152	17 Q57778	Q57778 pyrococcus
37	38	55.9	155	17 Q87LH7	Q87lh7 clostridium
38	38	55.9	203	16 Q8XLI7	Q8xli7 clostridium
39	38	55.9	222	17 Q97WC8	Q97wc8 sulfolobus
40	38	55.9	222	17 Q97WC8	Q97wc8 sulfolobus
41	38	55.9	313	5 Q7YTA2	Q7yta2 glomeris ma
42	38	55.9	325	16 Q9CBS2	Q9cbs2 mycobacteri
43	38	55.9	352	16 Q8G7R2	Q8g7r2 bifidobacte
44	38	55.9	353	17 Q8ZU64	Q8zu64 pyrobaculum
45	38	55.9	404	16 Q897K0	Q897k0 clostridium

## ALIGNMENTS

### RESULT 1

Q8K408 PRELIMINARY; PRT; 41 AA.  
ID Q8K408;  
AC Q8K408; (TRENBLrel. 22, Created)  
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)  
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)  
DE Truncated macrophage colony stimulating factor.  
GN CSPI.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
OX NCBI\_TaxID=10116;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=LEW tl;  
RX MEDLINE=22069908; PubMed=12074592;  
RA Dobbins D.E.; Sood R.; Hashimoto A.; Hansen C.T.; Wilder R.L.;  
RA Remmers E.F.;  
RT "Mutation of macrophage colony stimulating factor (Csfl) causes  
RT osteopetrosis in the tl rat.";  
RL Biochem Biophys. Res. Commun. 294:1114-1120 (2002).  
DR EMBL; AF514357; AAM54137.1;  
SQ SEQUENCE 41 AA; 4178 MW; ID342C19BD18AA41 CRC64;

### Query Match

Best Local Similarity 66.2%; Score 45; DB 11; Length 41;  
Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

### QY 1 PAVVGLSPGQEY 12

Db 22 PAAAGLSPEQE 33

### RESULT 2

Q9HPR8 PRELIMINARY; PRT; 821 AA.  
ID Q9HPR8;  
AC Q9HPR8;  
DT 01-MAR-2001 (TRENBLrel. 16, Created)

DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)  
 DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)  
 DE DNA helicase.  
 GN HEL OR VNG1501G.  
 OS Halobacterium sp. (strain NRC-1 / ATCC 700922 / JCM 11081).  
 OC Archaea; Euryarchaeota; Halobacteriales; Halobacteriaceae; Halobacterium.  
 OX NCBI\_TaxID=64091;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=20504483; PubMed=11016950;  
 RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,  
 RA Shukla H.D., Lasky S.R., Baliga N.S., Thorsson V., Sbrogna J.,  
 RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,  
 RA Leithauser B., Keller K., Cruz R., Danson M.J., Hough D.W.,  
 RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,  
 RA Isenbarger T.A., Peck R.F., Pohlshroder M., Spudis J.L., Jung K.-H.,  
 RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,  
 RA Ehardt H., Lowe T.M., Liang P., Riley M., Hood L., Dassarma S.,  
 RT "Genome sequence of Halobacterium species NRC-1."  
 RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).  
 DR EMBL; AE05064; AAC15799.1; --  
 DR PIR; C84304; C84304.  
 DR GO; GO:0004386; F:helicase activity; IEA.  
 DR GO; GO:0000166; F:nucleotide binding; IEA.  
 DR GO; GO:0007242; P:intracellular signaling cascade; IEA.  
 DR InterPro; IPR003593; AAA\_ATPase.  
 DR InterPro; IPR001478; PDZ.  
 DR SMART; SM00382; AAA; 1.  
 DR SMART; SM00228; PDZ; 1.  
 KW Helicase; Complete proteome.  
 SQ SEQUENCE 821 AA; 98848 MW; C454C76B984A5702 CRC64;  
  
 Query Match 63.2%; Score 43; DB 17; Length 821;  
 Best Local Similarity 90.0%; Pred. No. 91;  
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
 QY 2 AVVGLSPGEQ 11  
 DB 326 AVVGLSPGEQ 335  
  
 RESULT 3  
 ID Q9HKE3 PRELIMINARY; PRT; 228 AA.  
 AC Q9HKE3;  
 DT 01-MAR-2001 (TREMBlrel. 16, Created)  
 DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)  
 DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)  
 DE Precorrin-2 methyltransferase related protein.  
 GN TA0658.  
 OS Thermoplasma acidophilum.  
 OC Archaea; Euryarchaeota; Thermoplasma; Thermoplasmales;  
 OC Thermoplasmatocaeae; Thermoplasma.  
 OX NCBI\_TaxID=2303;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=20479972; PubMed=11029001;  
 RA Ruepp A., Graml W., Santos-Martinez M.-L., Koretke K.K., Volker C.,  
 RA Mewes H.-W., Frishman D., Stocker S., Lupas A.N., Baumeister W.;  
 RT "The genome sequence of the thermoacidophilic scavenger Thermoplasma  
 acidophilum."  
 RL Nature 407:508-513(2000).  
 DR EMBL; AL445065; CAC11796.1; --  
 DR GO; GO:0008757; F:S-adenosylmethionine-dependent methyltransf. . .; IEA.  
 DR GO; GO:0016740; F:transferase activity; IEA.  
 DR GO; GO:0008152; P:metabolism; IEA.  
 DR GO; GO:0006779; P:porphyrin biosynthesis; IEA.  
 DR InterPro; IPR006364; Cobi Cbil.  
 DR InterPro; IPR008078; Cor/por Metransf.  
 DR InterPro; IPR003043; Uropor Metransf.  
 DR Pfam; PF00590; TP\_methylase; 1.

DR TIGRFAMS; TIGR01467; cobi\_cbil; 1.  
 DR PROSITE; PS00839; SUMT\_1; 1.  
 KW Transferase; Methyltransferase; Complete proteome.  
 SQ SEQUENCE 228 AA; 25084 MW; 11ABD8B68192A67C CRC64;  
  
 Query Match 60.3%; Score 41; DB 17; Length 228;  
 Best Local Similarity 63.6%; Pred. No. 51;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
  
 QY 3 VVGLSPGEQEQ 13  
 DB 5 VVGLSPGEQDPD 15  
  
 RESULT 4  
 ID Q9UXP0 PRELIMINARY; PRT; 326 AA.  
 AC Q9UXP0;  
 DT 01-MAY-2000 (TREMBlrel. 13, Created)  
 DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)  
 DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)  
 DE F420-dependent N5,N10-methylene-tetrahydromethanopterin reductase,  
 DE putative.  
 GN PFDA.  
 OS Methanobolus tindarius.  
 OC Archaea; Euryarchaeota; Euryarchaeota orders incertae sedis;  
 OC Methanosarcinales; Methanosarcinaceae; Methanobolus.  
 OX NCBI\_TaxID=2221;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=DSM 2278;  
 RA Westenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,  
 RA Gottschalk G., Blaut M.;  
 RT "The F420H2-dehydrogenase from Methanobolus tindarius: Cloning of the  
 RT ffd operon and expression of the genes in Escherichia coli."  
 RL Submitted (SEP-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ011519; CAB56639.1; --  
 DR PIR; T45226; T45226.  
 DR InterPro; IPR002103; Bac\_luciferase.  
 DR Pfam; PF00296; bac\_luciferase; 1.  
 SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;  
  
 Query Match 60.3%; Score 41; DB 1; Length 326;  
 Best Local Similarity 70.0%; Pred. No. 75;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
  
 QY 2 AVVGLSPGEQ 11  
 DB 88 AILGLPGGEQ 97  
  
 RESULT 5  
 ID Q7X128 PRELIMINARY; PRT; 877 AA.  
 AC Q7X128;  
 DT 01-OCT-2003 (TREMBlrel. 25, Created)  
 DT 01-OCT-2003 (TREMBlrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)  
 DE VirB4.  
 GN VIRB4.  
 OS Xanthomonas campestris (pv. citri).  
 OG Plasmid pXCB.  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;  
 OC Xanthomonadaceae; Xanthomonas.  
 OX NCBI\_TaxID=346;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA Yuan Q., Brunings A.M., El-Yacoubi B., Shanker S., Gabriel D.W.;  
 RT "A self-mobilizing plasmid from a South American citrus canker strain  
 RT carries required pathogenicity gene pthB."  
 RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AY228335; AAC72100.1; --  
 KW Plasmid.

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SQ SEQUENCE 877 AA; 100012 MW; 83B94EDB23A0728D CRC64;
Query Match 60.3%; Score 41; DB 2; Length 877;
Best Local Similarity 63.6%; Pred. No. 2.2e+02;
Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 3 VVGLSPGQEQY 13
Db 785 VLGLTPGQYEF 795

RESULT 6
Q9L4W3 PRELIMINARY; PRT; 11096 AA.
AC Q9L4W3
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE NYSC.
GN Streptomyces noursei.
OS Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycinae; Streptomycetaceae; Streptomycetes.
OC Streptomycetaceae; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=1971;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 11455;
RX MEDLINE=20334850; PubMed=10873841;
RA Brattaset T., Sekurova O.N., Sletta H., Ellingsen T.E., Strom A.R.,
RA Valla S., Zetchev S.B.;
RT "Biosynthesis of the polyene antifungal antibiotic nystatin in
RT Streptomyces noursei ATCC 11455: analysis of the gene cluster and
RT deduction of the biosynthetic pathway.";
RL Chem. Biol. 7:395-403(2000).
DR EMBL; AF263912; AAF71776.1; -.
DR HSP; F25715; IMLA.
DR GO; GO:0004024; F:alcohol dehydrogenase activity, zinc-dependent; IEA.
DR GO; GO:0016740; F:transferase activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR GO; GO:0004314; F:[acyl-carrier protein] S-malonyltransferase. .; IEA.
DR GO; GO:0006633; F:fatty acid biosynthesis; IEA.
DR GO; GO:0008152; F:metabolism; IEA.
DR InterPro; IPR001227; Ac trans.
DR InterPro; IPR002085; Adh zn family.
DR InterPro; IPR004410; FAD.
DR InterPro; IPR000794; Ketoacyl synth.
DR InterPro; IPR006162; Ppantne S.
DR InterPro; IPR006163; Pp_bind.
DR Pfam; PF00698; Acyl_transf; 6.
DR Pfam; PF00107; ADH_zinc_N; 1.
DR Pfam; PF00109; ketoacyl-synt; 6.
DR Pfam; PF02801; ketoacyl-synt C; 6.
DR Pfam; PF00550; pp-binding; 6.
DR TIGRfam; TIGR00128; fadD; 6.
DR PROSITE; PS50075; ACP_DOMAIN; 6.
DR PROSITE; PS00606; B_KETOACYL_SYNTHASE; 6.
DR PROSITE; PS00012; PHOSPHOPANTHEINE; 5.
DR Phosphopantetheine; Transferase.
KW SEQUENCE 11096 AA; 1150415 MW; 776CAEAFCAE551DD CRC64;

Query Match 60.3%; Score 41; DB 2; Length 11096;
Best Local Similarity 63.6%; Pred. No. 3.4e+03;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGQEQ 11
Db 4998 PEVTLAPGDQ 5008

RESULT 7
Q97WD1 PRELIMINARY; PRT; 227 AA.
ID Q97WD1
AC Q97WD1
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE NYSC.
GN Streptomyces noursei.
OS Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycinae; Streptomycetaceae; Streptomycetes.
OC Streptomycetaceae; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=1971;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 11455;
RX MEDLINE=20334850; PubMed=10873841;
RA Brattaset T., Sekurova O.N., Sletta H., Ellingsen T.E., Strom A.R.,
RA Valla S., Zetchev S.B.;
RT "Biosynthesis of the polyene antifungal antibiotic nystatin in
RT Streptomyces noursei ATCC 11455: analysis of the gene cluster and
RT deduction of the biosynthetic pathway.";
RL Chem. Biol. 7:395-403(2000).
DR EMBL; AF263912; AAF71776.1; -.
DR HSP; F25715; IMLA.
DR GO; GO:0004024; F:alcohol dehydrogenase activity, zinc-dependent; IEA.
DR GO; GO:0016740; F:transferase activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR GO; GO:0004314; F:[acyl-carrier protein] S-malonyltransferase. .; IEA.
DR GO; GO:0006633; F:fatty acid biosynthesis; IEA.
DR GO; GO:0008152; F:metabolism; IEA.
DR InterPro; IPR001227; Ac trans.
DR InterPro; IPR002085; Adh zn family.
DR InterPro; IPR004410; FAD.
DR InterPro; IPR000794; Ketoacyl synth.
DR InterPro; IPR006162; Ppantne S.
DR InterPro; IPR006163; Pp_bind.
DR Pfam; PF00698; Acyl_transf; 6.
DR Pfam; PF00107; ADH_zinc_N; 1.
DR Pfam; PF00109; ketoacyl-synt; 6.
DR Pfam; PF02801; ketoacyl-synt C; 6.
DR Pfam; PF00550; pp-binding; 6.
DR TIGRfam; TIGR00128; fadD; 6.
DR PROSITE; PS50075; ACP_DOMAIN; 6.
DR PROSITE; PS00606; B_KETOACYL_SYNTHASE; 6.
DR PROSITE; PS00012; PHOSPHOPANTHEINE; 5.
DR Phosphopantetheine; Transferase.
KW SEQUENCE 11096 AA; 1150415 MW; 776CAEAFCAE551DD CRC64;

Query Match 60.3%; Score 41; DB 2; Length 11096;
Best Local Similarity 63.6%; Pred. No. 3.4e+03;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGQEQ 11
Db 4998 PEVTLAPGDQ 5008

RESULT 7
Q97WD1 PRELIMINARY; PRT; 227 AA.
ID Q97WD1
AC Q97WD1
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE NYSC.
GN Streptomyces noursei.
OS Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycinae; Streptomycetaceae; Streptomycetes.
OC Streptomycetaceae; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=1971;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 11455;
RX MEDLINE=20334850; PubMed=10873841;
RA Brattaset T., Sekurova O.N., Sletta H., Ellingsen T.E., Strom A.R.,
RA Valla S., Zetchev S.B.;
RT "Biosynthesis of the polyene antifungal antibiotic nystatin in
RT Streptomyces noursei ATCC 11455: analysis of the gene cluster and
RT deduction of the biosynthetic pathway.";
RL Chem. Biol. 7:395-403(2000).
DR EMBL; AF263912; AAF71776.1; -.
DR HSP; F25715; IMLA.
DR GO; GO:0004024; F:alcohol dehydrogenase activity, zinc-dependent; IEA.
DR GO; GO:0016740; F:transferase activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR GO; GO:0004314; F:[acyl-carrier protein] S-malonyltransferase. .; IEA.
DR GO; GO:0006633; F:fatty acid biosynthesis; IEA.
DR GO; GO:0008152; F:metabolism; IEA.
DR InterPro; IPR001227; Ac trans.
DR InterPro; IPR002085; Adh zn family.
DR InterPro; IPR004410; FAD.
DR InterPro; IPR000794; Ketoacyl synth.
DR InterPro; IPR006162; Ppantne S.
DR InterPro; IPR006163; Pp_bind.
DR Pfam; PF00698; Acyl_transf; 6.
DR Pfam; PF00107; ADH_zinc_N; 1.
DR Pfam; PF00109; ketoacyl-synt; 6.
DR Pfam; PF02801; ketoacyl-synt C; 6.
DR Pfam; PF00550; pp-binding; 6.
DR TIGRfam; TIGR00128; fadD; 6.
DR PROSITE; PS50075; ACP_DOMAIN; 6.
DR PROSITE; PS00606; B_KETOACYL_SYNTHASE; 6.
DR PROSITE; PS00012; PHOSPHOPANTHEINE; 5.
DR Phosphopantetheine; Transferase.
KW SEQUENCE 11096 AA; 1150415 MW; 776CAEAFCAE551DD CRC64;

Query Match 60.3%; Score 40; DB 17; Length 227;
Best Local Similarity 54.5%; Pred. No. 76;
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 VVGLSPGQEQY 13
Db 10 IVGVGPGDPEY 20

RESULT 8
Q7WCX1 PRELIMINARY; PRT; 242 AA.
ID Q7WCX1
AC Q7WCX1
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Bacteriophage-related DNA polymerase.
GN BB3808.
OS Bordetella bronchiseptica (Alcaligenes bronchisepticus).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Alcaligenaceae; Bordetella.
OX NCBI_TaxID=518;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=RB50 / ATCC BAA-588;
RX MEDLINE=22827954; PubMed=12910271;
RA Parkhill J., Sebaihia M., Preston A., Murphy L.D., Thomson N.,
RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,
RA Cerdano-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,
RA Achtman M., Atkin R., Baker S., Basham D., Bason N., Cherevach I.,
RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,
RA Felwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
RA Leather S., Moule S., Norberczak H., O'Neill S., Ormond D., Price C.,
RA Rabinowitch E., Rutter S., Sanders M., Saunders D., Seeger K.,
RA Sharp S., Simmonds M., Skelton J., Squares R., Stevens K.,
RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;
RT "Comparative analysis of the genome sequences of Bordetella pertussis,
RT Bordetella parapertussis and Bordetella bronchiseptica.";
RL Nat. Genet. 35:32-40(2003).
DR EMBL; BX640448; CAE35782.1; -.
KW Complete proteome.

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SQ SEQUENCE 242 AA; 25627 MW; EECAD9B319823284 CRC64;
  Query Match 58.8%; Score 40; DB 16; Length 242;
  Best Local Similarity 80.0%; Pred. No. 82;
  Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 VVGLSPGGEQ 12
DB 96 VVGEAPGGEQ 105

RESULT 9
Q7W5D6 PRELIMINARY; PRT; 298 AA.
AC Q7W5D6;
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DE Bacteriophage-related DNA polymerase.
GN BPP3357.
OS Bordetella parapertussis.
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Alcaligenaceae; Bordetella.
OX NCBI_TaxID=519;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=12822 / ATCC BAA-587;
RX MEDLINE=22827954; PubMed=12910271;
RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,
RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,
RA Cerdeno-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,
RA Achman M., Atkin R., Baker S., Basham D., Cronin A., Davis P., Doggett J.,
RA Chillingworth T., Collins M., Norberczak H., O'Neil S., Ormond D., Price C.,
RA Feltwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
RA Leather S., Moule S., Norberczak H., O'Neil S., Ormond D., Price C.,
RA Rabinowitsch E., Rutter S., Sanders M., Saunders D., Seeger K.,
RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,
RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;
RT "Comparative analysis of the genome sequences of Bordetella pertussis,
RT Bordetella parapertussis and Bordetella bronchiseptica.";
RL Nat. Genet. 35:32-40(2003).
DR EMBL; BX640433; CAE38642.1; -.
KW Complete proteome.
SQ SEQUENCE 298 AA; 31202 MW; A858C8199E2AC9B7 CRC64;

  Query Match 58.8%; Score 40; DB 16; Length 298;
  Best Local Similarity 80.0%; Pred. No. 1e+02;
  Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 VVGLSPGGEQ 12
DB 152 VVGEAPGGEQ 161

RESULT 10
Q7VTI2 PRELIMINARY; PRT; 298 AA.
AC Q7VTI2;
DT 01-OCT-2003 (TREMBLrel. 25, Created)
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Bacteriophage-related DNA polymerase.
GN BP3556.
OS Bordetella pertussis.
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Alcaligenaceae; Bordetella.
OX NCBI_TaxID=520;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Tohama I / ATCC BAA-589 / NCTC 13251;
RX MEDLINE=22827954; PubMed=12910271;
RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,
RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,

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RA Cerdeno-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,
RA Achman M., Atkin R., Baker S., Basham D., Cronin A., Davis P., Doggett J.,
RA Chillingworth T., Collins M., Norberczak H., O'Neil S., Ormond D., Price C.,
RA Feltwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
RA Leather S., Moule S., Norberczak H., O'Neil S., Ormond D., Price C.,
RA Rabinowitsch E., Rutter S., Sanders M., Saunders D., Seeger K.,
RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,
RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;
RT "Comparative analysis of the genome sequences of Bordetella pertussis,
RT Bordetella parapertussis and Bordetella bronchiseptica.";
RL Nat. Genet. 35:32-40(2003).
DR EMBL; BX640421; CAE43815.1; -.
KW Complete proteome.
SQ SEQUENCE 298 AA; 31231 MW; 9E4BC84B419506F3 CRC64;

  Query Match 58.8%; Score 40; DB 16; Length 298;
  Best Local Similarity 80.0%; Pred. No. 1e+02;
  Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 VVGLSPGGEQ 12
DB 152 VVGEAPGGEQ 161

RESULT 11
Q947A7 PRELIMINARY; PRT; 355 AA.
AC Q947A7;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Enolase (EC 4.2.1.11) (2-phosphoglycerate dehydratase) (2-phospho-D-
DE glycerate hydro-lyase) (Fragment).
GN ENO.
OS Nitellopsis obtusa.
OC Eukaryota; Viridiplantae; Streptophyta; Charales; Characeae;
OC Nitellopsis.
OX NCBI_TaxID=40811;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=X-854;
RX MEDLINE=21437986; PubMed=11526220;
RA Keeling P.J., Palmer J.D.;
RT "Lateral transfer at the gene and subgenic levels in the evolution of
RT eukaryotic enolase.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:10745-10750(2001).
CC -!- CATALYTIC ACTIVITY: 2-PHOSPHO-D-GLYCERATE = PHOSPHOENOLPYRUVATE +
CC H(2)O.
CC -!- COPACTOR: MAGNESIUM IS REQUIRED FOR CATALYSIS AND FOR STABILIZING
CC THE DIMER (BY SIMILARITY).
CC -!- PATHWAY: GLYCOLYSIS.
CC -!- SUBUNIT: HOMODIMER (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE ENOLASE FAMILY.
DR EMBL; AF348916; AAL05455.1; -.
DR GO; GO:0000015; C:phosphopyruvate hydratase complex; IEA.
DR GO; GO:0016829; F:lyase activity; IEA.
DR GO; GO:0000287; F:magnesium ion binding; IEA.
DR GO; GO:0004634; F:phosphopyruvate hydratase activity; IEA.
DR GO; GO:0006096; P:glycolysis; IEA.
DR InterPro; IPR000941; Enolase.
DR Pfam; PF00113; enolase; 1.
DR Pfam; PF03952; enolase.N; 1.
DR PRINTS; PR00148; ENOLASE.
DR ProDom; PD000902; Enolase; 1.
DR PROSITE; PS00164; ENOLASE; 1.
KW Glycolysis; Lyase; Magnesium.
FT NON_TER 1
FT NON_TER 355
SQ SEQUENCE 355 AA; 38295 MW; 400DF160087DE450 CRC64;

  Query Match 58.8%; Score 40; DB 10; Length 355;
  Best Local Similarity 54.5%; Pred. No. 1.2e+02;

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Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
QY 1 PAVVGLSPGEQ 11
Db 28 PAVIGMDPADQ 38

RESULT 12
Q8NHX1 PRELIMINARY; PRT; 357 AA.
AC Q8NHX1;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Extracellular signal-related kinase 1b.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Aebersold D.M., Yung Y., Seger R.;
RT "Properties of human ERK1b."
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY033607; AAK52329.1; -.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0004707; F:MAP kinase activity; IEA.
DR GO; GO:0004574; F:protein serine/threonine kinase activity; IEA.
DR GO; GO:0004713; F:protein-tyrosine kinase activity; IEA.
DR GO; GO:0016740; F:transferase activity; IEA.
DR GO; GO:0006468; P:protein amino acid phosphorylation; IEA.
DR InterPro; IPR008349; Erk 1 2 MAPK.
DR InterPro; IPR008350; Erk 3 4 MAPK.
DR InterPro; IPR003527; MAP Kin.
DR InterPro; IPR00719; Prot_kinase.
DR InterPro; IPR002290; Ser_Ehr_kinase.
DR InterPro; IPR001245; Tyr_pkinase.
DR Pfam; PF00069; pkinase; 1.
DR PRINTS; PR01770; ERK1ERK2MAPK.
DR PROSITE; PS01771; ERK3ERK4MAPK.
DR ProDom; PD000001; Prot_kinase; 1.
DR SMART; SM00220; S_TKC; 1.
DR SMART; SM00219; TYRK; 1.
DR PROSITE; PS01351; MAPK; 1.
DR PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
DR PROSITE; PS50011; PROTEIN_KINASE_DOM; 1.
KW ATP-binding; Kinase; Transferase
SQ SEQUENCE 357 AA; 4062 MW; 58C92773983ADA79 CRC64;

Query Match 58.8%; Score 40; DB 4; Length 357;
Best Local Similarity 72.7%; Pred. No. 1.2e+02;
Matches 8; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 11
Db 344 PAAVGLGAGEQ 354

RESULT 13
Q9NWD0 PRELIMINARY; PRT; 358 AA.
AC Q9NWD0;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Hypothetical protein.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Tissue=Embryo;

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RA Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
RA Nishikawa T., Nagai K., Sugano S., Shiratori A., Sudo H., Sugawara M.,
RA Wagatsuma M., Hosoi T., Kaku Y., Kodaira H., Kondo H., Sugawara M.,
RA Takahashi M., Chiba Y., Ishida S., Murakawa K., Ono Y., Takiguchi S.,
RA Watanabe S., Kimura K., Murakami K., Ishii S., Kawai Y., Saito K.,
RA Yamamoto J., Wakamatsu A., Nakamura Y., Nagahari K., Masuho Y.,
RA Ninomiya K., Iwayanagi T.;
RT "NEDO human cDNA sequencing project."
RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AK000977; BAA91452.1; -.
DR InterPro; IPR003892; CUE.
DR Pfam; PF02845; CUE; 1.
DR SMART; SM00546; CUE; 1.
KW Hypothetical protein.
SQ SEQUENCE 358 AA; 38760 MW; BCE1AA7E95C73BF0 CRC64;

Query Match 58.8%; Score 40; DB 4; Length 358;
Best Local Similarity 58.3%; Pred. No. 1.3e+02;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 12
Db 213 PAMAGPGPDQE 224

RESULT 14
Q9RRL5 PRELIMINARY; PRT; 381 AA.
AC Q9RRL5;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Cytochrome P450.
GN DR2473.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
RX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Farnham W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Utterback T., Zaleski C.,
RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RA "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1."
RL Science 286:1571-1577(1999).
CC -1- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.
DR EMBL; AE002076; AAF12016.1; -.
DR PIR; F75270; F75270.
DR TIGR; DR2473; -.
DR GO; GO:0004497; F:monooxygenase activity; IEA.
DR GO; GO:0006118; F:electron transport; IEA.
DR InterPro; IPR001128; Cytochrome_P450.
DR Pfam; PF00067; P450; 1.
DR PRINTS; PR00385; P450.
DR PROSITE; PS00086; CYTOCHROME_P450; 1.
KW Heme; Monooxygenase; Oxidoreductase; Complete proteome.
SQ SEQUENCE 381 AA; 41940 MW; F191EA69F179B53 CRC64;

Query Match 58.8%; Score 40; DB 16; Length 381;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PAVVGLSP 8
Db 53 PAVVGLSP 60

```

RESULT 15  
Q9NWM3 PRELIMINARY; PRT; 386 AA.  
ID Q9NWM3  
AC Q9NWM3;  
DT 01-OCT-2000 (TrEMBLrel. 15, Created)  
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)  
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
DE Hypothetical protein FLJ20739.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
RP SEQUENCE FROM N.A.  
RA Kawakami T., Noguchi S., Itoh T., Shigeta K., Senba T., Matsumura K.,  
RA Nakajima Y., Mizuno T., Morinaga M., Ota T., Suzuki Y., Obayashi M.,  
RA Nishi T., Shibahara T., Tanaka T., Nakamura Y., Isegai T., Sugano S.;  
RT "NEDO human cDNA sequencing project.";  
RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.  
DR EMBL: AK000746; BAA91357.1; -  
DR Genew; HGNC:19843; CL4orf34.  
DR InterPro; IPR003892; CUE.  
DR Pfam; PF02845; CUE; 1.  
DR SMART; SM00546; CUE; 1.  
KW Hypothetical protein.  
SQ SEQUENCE 386 AA; 42258 MW; F5530FA47C267895 CRC64;  
Query Match 58.8%; Score 40; DB 4; Length 386;  
Best Local Similarity 58.3%; Pred. No. 1.4e+02;  
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;  
QY 1 PAVVGLSPGQOE 12  
Db 213 PAVAGGPGDQOE 224  
Search completed: May 7, 2004, 12:37:55  
Job time : 25.4767 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 36.27 Seconds  
(without alignments)  
101.272 Million cell updates/sec

Title: US-09-786-214A-13  
Perfect score: 68  
Sequence: 1 PAVVGLSPGEQY 13

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues  
Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : A Genesep 29Jan04:\*  
1: Genesep1980s:\*  
2: Genesep1930s:\*  
3: Genesep2000s:\*  
4: Genesep2001s:\*  
5: Genesep2002s:\*  
6: Genesep2003as:\*  
7: Genesep2003bs:\*  
8: Genesep2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	68	100.0	13	3 AAY84267	Aay84267 Peptide d
2	68	100.0	14	3 AAY84266	Aay84266 Peptide d
3	68	100.0	20	3 AAY84265	Aay84265 Truncated
4	68	100.0	25	3 AAY84264	Aay84264 Peptide o
5	61	89.7	13	3 AAY84268	Aay84268 Peptide d
6	61	89.7	15	3 AAY84269	Aay84269 Peptide d
7	45	66.2	234	4 AAB36208	Ab36208 Human imm
8	41	60.3	106	5 ABJ10397	Abj10397 Mutant an
9	41	60.3	106	5 ABJ10395	Abj10395 Mutant an
10	41	60.3	11096	4 AAE10129	Aae10129 Streptomy
11	40	58.8	358	4 AAB92530	Aab92530 Human pro
12	40	58.8	359	5 ABB97563	Abb97563 Novel hum
13	40	58.8	668	3 AAB57055	Aab57055 Human pro
14	39	57.4	18	2 AAY41875	Aay41875 Rheumatoi
15	39	57.4	18	4 AAU25388	Aau25388 Schizophr
16	39	57.4	18	4 AAU15732	Aau15732 Schizophr
17	39	57.4	18	5 ABG78871	Abg78871 Multiple
18	39	57.4	23	5 ABP62618	Abp62618 Human imm
19	39	57.4	74	6 ABU18694	Abu18694 Antibody
20	39	57.4	88	3 AAY56655	Aay56655 Human p
21	39	57.4	93	5 AAU08983	Aau08983 Human ant
22	39	57.4	94	7 ADD69248	Add69248 Human lig
23	39	57.4	95	6 ABO27154	Abo27154 Human ger
24	39	57.4	95	6 ABO27153	Abo27153 Human ger
25	39	57.4	96	6 ABO27155	Abo27155 Human ger

26	39	57.4	96	6 ABO27150	Abo27150 Human ger
27	39	57.4	100	5 AAE23987	Aae23987 Human MOC
28	39	57.4	104	2 AAW31123	Aaw31123 Alpha lig
29	39	57.4	104	2 AAW26795	Aaw26795 Anti-gp54
30	39	57.4	105	7 ABR61567	Abr61567 HIV-1 neu
31	39	57.4	106	2 AAR41234	Aar41234 Monoclonona
32	39	57.4	106	2 AAW71241	Aaw71241 Light cha
33	39	57.4	106	6 AAE34874	Aae34874 BIWA4 ant
34	39	57.4	106	6 AAE34875	Aae34875 BIWA4 ant
35	39	57.4	106	6 AAE33419	Aae33419 Murine KS
36	39	57.4	106	6 AAE33421	Aae33421 Murine KS
37	39	57.4	106	6 AAO30909	Aao30909 hu-KS ant
38	39	57.4	107	2 AAR32129	Aar32129 Anti-TL2R
39	39	57.4	107	2 AAR37610	Aar37610 B-R10 MAB
40	39	57.4	107	2 AAR50190	Aar50190 Light cha
41	39	57.4	107	2 AAW08949	Aaw08949 Kappa lig
42	39	57.4	107	2 AAW08948	Aaw08948 Kappa lig
43	39	57.4	107	2 AAW24514	Aaw24514 Anti-Fact
44	39	57.4	107	2 AAW24515	Aaw24515 Anti-Fact
45	39	57.4	107	2 AAW24513	Aaw24513 Anti-Fact

ALIGNMENTS

RESULT 1  
AAY84267  
ID AAY84267 standard; peptide; 13 AA.  
XX  
AC AAY84267;  
XX  
XX 12-JUL-2000 (first entry)  
XX  
DE Peptide derived from macrophage colony stimulating gene alternative ORF.  
XX  
KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
XX  
OS Synthetic.  
OS Homo sapiens.  
XX  
PN W0200013699-Al.  
XX  
PD 16-MAR-2000.  
XX  
PF 03-SEP-1999; 99WO-US020344.  
XX  
PR 04-SEP-1998; 98US-0099077P.  
XX  
PA (LUDW-) LUDWIG INST CANCER RES.  
XX  
PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
XX  
DR WPI; 2000-256859/22.  
XX  
PT Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.  
XX  
PS Example 2; Page 40; 74pp; English.

The present sequence represents a peptide which is derived from a tumour rejection antigen precursor encoded by an alternative open reading frame (ORF) of human macrophage colony stimulating gene. Peptides derived from the alternative ORF of macrophage-colony stimulating factor, when presented by an antigen presenting cell having a human leukocyte antigen (HLA) class I molecule, effectively induce the activation and proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF of macrophage-colony stimulating factor are useful for enriching selectively a population of T lymphocytes with CD8+ T lymphocytes. They are also useful for diagnosing a disorder characterized by expression of the polypeptide, and for identifying

CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 100.0%; Score 68; DB 3; Length 13;  
Best Local Similarity 100.0%; Pred. No. 0.00015; Mismatches 0; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQY 13  
| | | | | | | | | |  
Db 1 PAVVGLSPGEQY 13

RESULT 2

AY84266  
ID AAY84266 standard; peptide; 14 AA.

AC AAY84266;

DT 12-JUL-2000 (first entry)

Peptide derived from macrophage colony stimulating gene alternative ORF.

KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Synthetic.

OS Homo sapiens.

PN WO200013699-A1.

PD 16-MAR-2000.

PF 03-SEP-1999; 99WO-US020344.

PR 04-SEP-1998; 98US-0099077P.

PA (LUDW-) LUDWIG INST CANCER RES.

PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

DR WPI; 2000-256859/22.

PT Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Claim 2; Page 39; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 14 AA;

Query Match 100.0%; Score 68; DB 3; Length 14;  
Best Local Similarity 100.0%; Pred. No. 0.00016; Mismatches 0; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQY 13  
| | | | | | | | | |  
Db 2 PAVVGLSPGEQY 14

RESULT 3

AY84265

ID AAY84265 standard; peptide; 20 AA.

AC AAY84265;

DT 12-JUL-2000 (first entry)

Truncated macrophage colony stimulating factor tumour antigen.

KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Homo sapiens.

PN WO200013699-A1.

PD 16-MAR-2000.

PF 03-SEP-1999; 99WO-US020344.

PR 04-SEP-1998; 98US-0099077P.

PA (LUDW-) LUDWIG INST CANCER RES.

PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

DR WPI; 2000-256859/22.

DR N-PSDB; AAZ99675.

PT Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Claim 3; Page 64; 74pp; English.

XX The present sequence represents a truncated tumour rejection antigen  
CC precursor, and is encoded by a truncated alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 20 AA;

Query Match 100.0%; Score 68; DB 3; Length 20;  
Best Local Similarity 100.0%; Pred. No. 0.00024;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQY 13  
| | | | | | | | | |  
Db 5 PAVVGLSPGEQY 17

RESULT 4

AY84264

ID AAY84264 standard; peptide; 25 AA.

AC AAY84264;

DT 12-JUL-2000 (first entry)

DE Peptide of alternate reading frame of macrophage colony stimulating gene.  
XX Renal cell carcinoma; antigen; cytotoxic T lymphocyte;

KW tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Homo sapiens.

PN WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX N-PSDB; AAZ99672.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 1; Page 64; 74pp; English.

XX The present sequence represents a tumour rejection antigen precursor, and  
 CC is encoded by an alternative open reading frame (ORF) of human macrophage  
 CC colony stimulating gene. Peptides derived from the alternative ORF of  
 CC macrophage-colony stimulating factor, when presented by an antigen  
 CC presenting cell having a human leukocyte antigen (HLA) class I molecule,  
 CC effectively induce the activation and proliferation of CD8+ cytotoxic T  
 CC lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF  
 CC of macrophage-colony stimulating factor are useful for enriching  
 CC selectively a population of T lymphocytes with CD8+ T lymphocytes. They  
 CC are also useful for diagnosing a disorder characterized by expression of  
 CC the polypeptide, and for identifying functional variants and mimetics

XX Sequence 25 AA;

Query Match 100.0%; Score 68; DB 3; Length 25;

Best Local Similarity 100.0%; Pred. NO. 0.00031;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 PAVVGLSPGGEY 13

Db 5 PAVVGLSPGGEY 17

RESULT 5

AAV84268

ID AAV84268 standard; peptide; 13 AA.

XX AAV84268;

XX 12-JUL-2000 (first entry)

XX Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.

OS Homo sapiens.

XX WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

PR 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 89.7%; Score 61; DB 3; Length 13;

Best Local Similarity 100.0%; Pred. NO. 0.0022;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 PAVVGLSPGGEY 12

Db 2 PAVVGLSPGGEY 13

RESULT 6

AAV84269

ID AAV84269 standard; peptide; 15 AA.

XX AAV84269;

XX 12-JUL-2000 (first entry)

XX Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.

OS Homo sapiens.

XX WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics  
 XX  
 SQ Sequence 15 AA;

Query Match 89.7%; Score 61; DB 3; Length 15;  
 Best Local Similarity 100.0%; Pred. No. 0.0026;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 12  
 DB 4 PAVVGLSPGEQ 15

RESULT 7  
 AAB36208  
 ID AAB36208 standard; protein; 234 AA.

AC AAB36208;

DT 15-FEB-2001 (first entry)

DE Human immune system associated protein HISAP-6.

KW Human; immune system associated protein; HISAP-6; immune disorder;  
 infection; autoimmune disease; cancer.

OS Homo sapiens.

PN US6135941-A.

PD 24-OCT-2000.

PF 27-MAR-1998; 98US-00049672.

PR 27-MAR-1998; 98US-00049672.

PA (INCY-) INCYTE PHARM INC.

PI Tang YT, Yue H, Lal P, Corley NC, Guegler KJ, Baughn MR;

PI Hillman JL, Au-Young J;

DR WPI: 2001-030926/04.

DR N-PSDB; AAC66524.

XX New human immune system associated proteins (HISAP) and polynucleotides  
 PT encoding the HISAP, useful for diagnosing, treating or preventing immune  
 PT or cell proliferative disorders or infections.

PS Claim 1; Col 59-60; 54pp; English.

XX The present invention provides the coding and protein sequences for a  
 CC number of human immune system associated proteins (HISAPs). These can be  
 CC used in the diagnosis and treatment of various autoimmune disorders,  
 CC infections and cell proliferation diseases. The diseases include AIDS,  
 CC adult respiratory distress syndrome, anaemia, asthma, atherosclerosis,  
 CC Crohn's disease, irritable bowel syndrome, multiple sclerosis, myasthenia  
 CC gravis, osteoarthritis, rheumatoid arthritis, scleroderma, systemic lupus  
 CC erythematosus, arteriosclerosis, cirrhosis and cancer  
 XX  
 SQ Sequence 234 AA;

Query Match 66.2%; Score 45; DB 4; Length 234;  
 Best Local Similarity 72.7%; Pred. No. 25;  
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 11  
 DB 28 PAIVLSLSPGER 38

RESULT 8

ABJ10397  
 ID ABJ10397 standard; protein; 106 AA.

AC ABJ10397;

DT 28-NOV-2002 (first entry)

DE Mutant anti-mesothelin Fv (ST6) variable light chain.

KW Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.

OS Unidentified.

OS Synthetic.

PN WO200240545-A2.

PD 23-MAY-2002.

PF 16-NOV-2001; 2001WO-US043602.

PR 17-NOV-2000; 2000US-0249805P.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Pastan IH, Onda M, Nagata S, Tsutsumi Y, Vincent JJ, Kreitman RJ;

PI Vasmatazis G, Lee B;

XX WPI: 2002-500208/53.

DR N-PSDB; APT08085.

XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.

PS Claim 12; Fig 2; 82pp; English.

XX The invention comprises the amino acid and coding sequences of  
 CC recombinant immunotoxin proteins. The immunotoxin proteins of the  
 CC invention contain an antibody (or an antigen-binding fragment) with a  
 CC substitution of a negatively charged amino acid for an uncharged or  
 CC positively charged amino acid. The immunotoxins of the invention have  
 CC reduced liver toxicity. The immunotoxins of the invention are useful for  
 CC killing a malignant cell (e.g. a cancer cell). The present amino acid  
 CC sequence represents a recombinant immunotoxin of the invention  
 XX

SQ Sequence 106 AA;

Query Match 60.3%; Score 41; DB 5; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 49;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 PAVVGLSPGEQ 11  
 DB 8 PAIVLSLSPGER 18

RESULT 9

ABJ10395  
 ID ABJ10395 standard; protein; 106 AA.

XX AC ABJ10395;







XX DT 27-JUN-2002 (first entry)  
 XX DE Novel human protein SEQ ID NO: 831.  
 XX KW Human; antianaemic; vulnary; antiinflammatory; immunomodulator;  
 KW antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy;  
 KW neuroprotective; antiparkinsonian; protein therapy; EST;  
 XX expressed sequence tag.  
 XX OS Homo sapiens.  
 XX PN WO200222660-A2.  
 XX PD 21-MAR-2002.  
 XX PF 10-SEP-2001; 2001WO-US026015.  
 XX PR 11-SEP-2000; 2000US-00659671.  
 XX PA (HYSE-) HYSEQ INC.  
 XX PI Tang YT, Liu C, Zhou P, Asundi V, Zhang J, Zhao QA, Ren F;  
 PI Xue AJ, Yang Y, Wehrman T, Drmanac RT;  
 DR WPI; 2002-292408/33.  
 DR N-PSDB; AAB32749.  
 XX An isolated polynucleotide for treating diseases associated with its  
 PT encoded polypeptide such as cancer and multiple sclerosis.  
 XX Claim 20; SEQ ID NO 831; 509pp; English.  
 CC The present invention provides the protein and coding sequences of 444  
 CC novel human proteins. These were isolated from expressed sequences tags  
 CC (ESTs). They can be used to stimulate cell growth, to regulate  
 CC haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth  
 CC e.g. in burn treatment, to regulate the immune system e.g. to treat  
 CC multiple sclerosis, to regulate activin or inhibin e.g. to treat  
 CC infertility, to regulate haemostasis or thrombolysis e.g. to treat stroke  
 CC and cancer, to screen for drugs, to treat inflammatory conditions e.g.  
 CC rheumatoid arthritis, and to treat nervous system disorders e.g.  
 CC Parkinson's disease. The present sequence is a protein of the invention  
 XX Sequence 359 AA;  
 SQ Query Match 58.8%; Score 40; DB 5; Length 359;  
 Best Local Similarity 58.3%; Pred. No. 2.8e+02;  
 Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;  
 QY 1 PAVVGLSPGQGE 12  
 ||: | ||: ||  
 Db 213 PAVAGPGGQGE 224  
 RESULT 13  
 AAB57055  
 ID AAB57055 standard; protein; 668 AA.  
 XX AAB57055;  
 XX 13-MAR-2001 (first entry)  
 DE Human prostate cancer antigen protein sequence SEQ ID NO:1633.  
 XX Human; prostate cancer; prostate cancer antigen; detection; diagnosis;  
 KW neuroprotective; cytostatic; cardioactive; immunomodulatory; muscular;  
 KW vulnary; gastrointestinal; nephrotropic; antiinfective; gynaecological;  
 KW antibacterial; gene therapy; neural; immune; reproductive; renal;  
 KW gastrointestinal; pulmonary; cardiovascular; proliferative disorder;  
 KW wound; infectious disease.  
 XX OS Homo sapiens.

XX FN WO2000055174-A1.  
 XX PD 21-SEP-2000.  
 XX PF 08-MAR-2000; 2000WO-US005988.  
 XX PR 12-MAR-1999; 99US-0124270P.  
 XX PA (HUMA-) HUMAN GENOME SCI INC.  
 PA (ROSE/) ROSEN C A.  
 XX PI Rosen CA, Ruben SM;  
 XX WPI; 2000-587513/55.  
 DR N-PSDB; AAF16258.  
 XX Prostate cancer associated gene sequences, referred to as prostate cancer  
 PT antigens, useful for treatment, prevention, and diagnosis of disorders  
 PT such as prostate cancer.  
 XX Claim 11; Page 2091-2094; 2338pp; English.  
 XX AAF15566 to AAF16505 encode the human prostate cancer associated  
 CC proteins, called prostate cancer antigens, given in AAB56363 to AAB57302.  
 CC The prostate cancer antigens can have neuroprotective, cytostatic,  
 CC cardioactive, immunomodulatory, muscular, vulnary, gastrointestinal,  
 CC nephrotropic, antiinfective, gynaecological and antibacterial activities,  
 CC and can be used in gene therapy. The prostate cancer antigen  
 CC polynucleotides may be used for detection of prostate cancer, chromosome  
 CC identification, as chromosome markers, and for numerous other diagnostic  
 CC or research purposes. The prostate cancer antigens may be used to treat  
 CC disorders such as neural, immune, muscular, reproductive,  
 CC gastrointestinal, pulmonary, cardiovascular, renal, and proliferative  
 CC disorders, wounds, and infectious diseases. AAF16506 to AAF16514 to  
 CC AAB57303 represent sequences used in the exemplification of the present  
 CC invention  
 XX SQ Sequence 668 AA;  
 Query Match 58.8%; Score 40; DB 3; Length 668;  
 Best Local Similarity 72.7%; Pred. No. 5.5e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 3 VVGLSPGQGEY 13  
 |||||:  
 Db 27 VVGLSPGQGYF 37  
 RESULT 14  
 AAY41875  
 ID AAY41875 standard; peptide; 18 AA.  
 XX AAY41875;  
 XX 09-DEC-1999 (first entry)  
 DE Rheumatoid arthritis diagnostic protein isoform peptide #26.  
 XX Human; rheumatoid arthritis; RA; diagnosis; RPI; RADF; detection;  
 KW rheumatoid arthritis diagnostic feature; ERPI; synovial fluid;  
 KW rheumatoid arthritis diagnostic protein isoform; screening;  
 KW expression reference protein isoform; prognosis.  
 XX OS Homo sapiens.  
 XX WO9947925-A2.  
 XX 23-SEP-1999.  
 XX 15-MAR-1999; 99WO-GB000763.  
 XX 13-MAR-1998; 98GB-00005477.

XX (OXFO-) OXFORD GLYCOSCIENCES UK LTD.  
 XX Parekh RB, Patel TP, Townsend RR;  
 XX WPI; 1999-571871/48.  
 DR Diagnosis of human rheumatoid arthritis by two-dimensional  
 PT electrophoresis.  
 XX Disclosure; Page 18; 157pp; English.  
 CC A method has been developed for the diagnosis of human rheumatoid  
 CC arthritis (RA) using two-dimensional electrophoresis to generate a two-  
 CC dimensional array of features. The method can be used for screening,  
 CC diagnosis and prognosis of RA in a subject or for monitoring the effect  
 CC of an anti-RA drug or therapy administered to a subject. The method  
 CC comprises: (a) analysing a sample of serum or plasma and optionally  
 CC synovial fluid by two-dimensional electrophoresis, to generate a two-  
 CC dimensional array of features; (b) identifying at least one chosen  
 CC feature whose relative abundance correlates with the presence or absence  
 CC of RA; and (c) comparing the abundance of each chosen feature in the  
 CC sample with the abundance of that chosen feature in serum or plasma from  
 CC one or more persons without RA, where the relative abundance of the  
 CC chosen feature or features in the sample indicates the presence or  
 CC absence of RA in the subject. The method can also be used in clinical  
 CC studies for testing drugs for therapy of RA, for purification of RA-  
 CC diagnostic protein isoforms (RPIs), and for production of antibodies to  
 CC RPIs. The RA-diagnostic feature (RADF) proteins can be used to identify  
 CC compounds that promote or inhibit their activity, which are then used as  
 CC RA drugs. Nucleic acid encoding RADFs can be used in gene therapy  
 CC protocols. AA41844 to AA42100 represent RPI peptides, AA42101 to  
 CC AA42103 represent expression reference protein isoform peptides and  
 CC AA225066 to AA225068 represent degenerate probes for RPIs, which are all  
 CC used in the exemplification of the present invention  
 XX Sequence 18 AA;  
 SQ Query Match 57.4%; Score 39; DB 2; Length 18;  
 Best Local Similarity 63.6%; Pred. No. 15;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 PAVVGLSPGEG 11  
 |||:|||||:  
 Db 8 PATLSLSPGER 18  
 Search completed: May 7, 2004, 12:33:46  
 Job time : 37.27 secs

PI Herath HMC, Parekh RB, Rohlf C, Terrett JA, Tyson KL;  
 XX WPI; 2001-570624/64.  
 DR New schizophrenia associated protein isoforms and encoding nucleic acid  
 PT molecules, useful for treatment, diagnosis and prognosis of schizophrenia  
 PT and screening for potential drugs for treatment and new drug targets.  
 XX Disclosure; Page 41; 148pp; English.  
 CC The sequence represents a schizophrenia-associated protein isoform (SPI).  
 CC These protein isoforms, e.g. SPI-206, SPI-238 and SPI-240 are detectable  
 CC in cerebrospinal fluid, serum or plasma and are useful markers of  
 CC schizophrenia. The sequences can be used for treatment and diagnosis of  
 CC schizophrenia, screening, prognosis, monitoring the results of therapy,  
 CC identifying patients most likely to respond to a particular therapy and  
 CC identification of new targets for drug treatment. SPI DNA is useful as a  
 CC nucleic acid probe to detect the presence of nucleic acids or SPIs  
 XX Sequence 18 AA;  
 SQ Query Match 57.4%; Score 39; DB 4; Length 18;  
 Best Local Similarity 63.6%; Pred. No. 15;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 PAVVGLSPGEG 11  
 |||:|||||:  
 Db 8 PATLSLSPGER 18  
 Search completed: May 7, 2004, 12:33:46  
 Job time : 37.27 secs

XX (OXFO-) OXFORD GLYCOSCIENCES UK LTD.  
 XX Parekh RB, Patel TP, Townsend RR;  
 XX WPI; 1999-571871/48.  
 DR Diagnosis of human rheumatoid arthritis by two-dimensional  
 PT electrophoresis.  
 XX Disclosure; Page 18; 157pp; English.  
 CC A method has been developed for the diagnosis of human rheumatoid  
 CC arthritis (RA) using two-dimensional electrophoresis to generate a two-  
 CC dimensional array of features. The method can be used for screening,  
 CC diagnosis and prognosis of RA in a subject or for monitoring the effect  
 CC of an anti-RA drug or therapy administered to a subject. The method  
 CC comprises: (a) analysing a sample of serum or plasma and optionally  
 CC synovial fluid by two-dimensional electrophoresis, to generate a two-  
 CC dimensional array of features; (b) identifying at least one chosen  
 CC feature whose relative abundance correlates with the presence or absence  
 CC of RA; and (c) comparing the abundance of each chosen feature in the  
 CC sample with the abundance of that chosen feature in serum or plasma from  
 CC one or more persons without RA, where the relative abundance of the  
 CC chosen feature or features in the sample indicates the presence or  
 CC absence of RA in the subject. The method can also be used in clinical  
 CC studies for testing drugs for therapy of RA, for purification of RA-  
 CC diagnostic protein isoforms (RPIs), and for production of antibodies to  
 CC RPIs. The RA-diagnostic feature (RADF) proteins can be used to identify  
 CC compounds that promote or inhibit their activity, which are then used as  
 CC RA drugs. Nucleic acid encoding RADFs can be used in gene therapy  
 CC protocols. AA41844 to AA42100 represent RPI peptides, AA42101 to  
 CC AA42103 represent expression reference protein isoform peptides and  
 CC AA225066 to AA225068 represent degenerate probes for RPIs, which are all  
 CC used in the exemplification of the present invention  
 XX Sequence 18 AA;  
 SQ Query Match 57.4%; Score 39; DB 2; Length 18;  
 Best Local Similarity 63.6%; Pred. No. 15;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 PAVVGLSPGEG 11  
 |||:|||||:  
 Db 8 PATLSLSPGER 18  
 Search completed: May 7, 2004, 12:33:46  
 Job time : 37.27 secs

RESULT 15  
 AAU25388  
 ID AAU25388 standard; peptide; 18 AA.  
 XX AAU25388;  
 AC  
 DT 18-DEC-2001 (first entry)  
 XX  
 DE Schizophrenia-Associated Protein Isoform (SPI) peptide #617.  
 XX  
 XX Schizophrenia-associated protein isoform; SPI; SPI-206; SPI-238; SPI-240;  
 KW neuroleptic; gene therapy; cerebrospinal fluid; serum; plasma.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200162785-A2.  
 XX  
 PD 30-AUG-2001.  
 XX  
 PF 23-FEB-2001; 2001WO-GB0000792.  
 XX  
 PR 24-FEB-2000; 2000GB-00004415.  
 PR 28-DEC-2000; 2000US-00750395.  
 XX  
 PA (OXFO-) OXFORD GLYCOSCIENCES UK LTD.  
 XX

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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:26:40 ; Search time 24.31 Seconds  
(without alignments)

168.726 Million cell updates/sec

Title: US-09-786-214A-14

Perfect score: 65

Sequence: 1 LPAVVGSLSPGEQ 13

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: sp\_archea.\*  
2: sp\_bacteria.\*  
3: sp\_fungi.\*  
4: sp\_human.\*  
5: sp\_invertebrate.\*  
6: sp\_mammal.\*  
7: sp\_mhc.\*  
8: sp\_organelle.\*  
9: sp\_phase.\*  
10: sp\_plant.\*  
11: sp\_rodent.\*  
12: sp\_virus.\*  
13: sp\_vertebrate.\*  
14: sp\_unclassified.\*  
15: sp\_rvirs.\*  
16: sp\_bacteriap.\*  
17: sp\_archaeap.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	49	75.4	41	11 Q8K408	Q8K408 rattus norv
2	44	67.7	381	16 Q9RRL5	Q9RRL5 deinococcus
3	43.5	66.9	614	16 Q8Y280	Q8Y280 raiostonia s
4	43	66.2	821	17 Q9HPR8	Q9HPR8 halobacteri
5	42	64.6	1541	6 Q8HX13	Q8HX13 sus scrofa
6	41	63.1	326	1 Q9UXP0	Q9UXP0 methanolobu
7	41	63.1	579	4 Q8N158	Q8N158 homo sapien
8	41	63.1	11096	2 Q9L4W3	Q9L4W3 streptomyce
9	40	61.5	154	17 Q8ZYJ3	Q8ZYJ3 pyrobaculum
10	40	61.5	242	16 Q7WCX1	Q7WCX1 bordetella
11	40	61.5	261	16 Q8NOA2	Q8NOA2 corynebacte
12	40	61.5	298	16 Q7WSD6	Q7WSD6 bordetella
13	40	61.5	298	16 Q7VTI2	Q7VTI2 bordetella
14	40	61.5	355	10 Q947A7	Q947A7 nitellopsis
15	40	61.5	357	4 Q8NHX1	Q8NHX1 homo sapien
16	40	61.5	358	4 Q9NWD0	Q9NWD0 homo sapien

17	40	61.5	386	4 Q9NWM3	Q9NWM3 homo sapien
18	40	61.5	428	16 Q9ZAG9	Q9ZAG9 listeria in
19	40	61.5	428	16 Q8Y661	Q8Y661 listeria mo
20	40	61.5	540	16 Q9RR71	Q9RR71 deinococcus
21	40	61.5	753	16 Q89T31	Q89T31 bradyrhizob
22	39	60.0	197	17 Q8PW19	Q8PW19 methanosarc
23	39	60.0	209	17 Q8TWT1	Q8TWT1 methanosarc
24	39	60.0	209	17 Q8TH81	Q8TH81 methanosarc
25	39	60.0	319	16 Q8ZKQ5	Q8ZKQ5 salmonella
26	39	60.0	319	16 Q8Z2X4	Q8Z2X4 salmonella
27	39	60.0	407	2 Q9LCW0	Q9LCW0 streptomyce
28	39	60.0	626	16 Q7WVC3	Q7WVC3 bordetella
29	39	60.0	627	16 Q7WH05	Q7WH05 bordetella
30	39	60.0	627	16 Q7W9Q3	Q7W9Q3 bordetella
31	39	60.0	1400	11 Q7TME1	Q7TME1 mus musculu
32	38.5	59.2	656	16 Q9RW14	Q9RW14 deinococcus
33	38	58.5	62	16 Q7VER3	Q7VER3 mycobacteri
34	38	58.5	136	11 Q61061	Q61061 mus musculu
35	38	58.5	152	17 Q9V2S2	Q9V2S2 pyrococcus
36	38	58.5	152	17 Q8TZK1	Q8TZK1 pyrococcus
37	38	58.5	155	17 Q57778	Q57778 pyrococcus
38	38	58.5	177	16 Q8U7R5	Q8U7R5 agrobacteri
39	38	58.5	212	5 Q9NA60	Q9NA60 caenorhabdi
40	38	58.5	222	17 Q97WC8	Q97WC8 sulfolobus
41	38	58.5	313	5 Q7YTA2	Q7YTA2 glomeris ma
42	38	58.5	325	16 Q9CBS2	Q9CBS2 mycobacteri
43	38	58.5	335	16 Q8ZKR7	Q8ZKR7 streptomyce
44	38	58.5	352	16 Q8G7R2	Q8G7R2 bifidobacte
45	38	58.5	383	16 Q7V8U3	Q7V8U3 prochloroco

## ALIGNMENTS

### RESULT 1

Q8K408 PRELIMINARY; PRT; 41 AA.  
AC Q8K408; 22 (TREMBLrel. 22, Created)  
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)  
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)  
DE Truncated macrophage colony stimulating factor.  
GN CSF1.  
OS Rattus norvegicus (Rat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
OX NCBI\_TaxID=10116;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=Lew tl;  
RX MEDLINE=22069908; PubMed=12074592;  
RA Dobbins D.E.; Sood R.; Hashiramoto A.; Hansen C.T.; Wilder R.L.;  
RA Remmers E.F.;  
RT "Mutation of macrophage colony stimulating factor (Csf1) causes  
RT osteopetrosis in the tl rat."  
RL Biochem. Biophys. Res. Commun. 294:1114-1120(2002).  
DR EMBL; AF514357; AAM54137.1; -  
SQ SEQUENCE 41 AA; 4178 MW; ID342C19BD18AA41 CRC64;

Query Match 75.4%; Score 49; DB 11; Length 41;  
Best Local Similarity 76.9%; Pred. No. 0.44;  
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 LPAVVGSLSPGEQ 13  
||| ||||| |||  
Db 21 LPAAGLSPREQE 33

### RESULT 2

Q9RRL5 PRELIMINARY; PRT; 381 AA.  
ID Q9RRL5  
AC Q9RRL5;  
DT 01-MAY-2000 (TREMBLrel. 13, Created)

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DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Cytochrome P450.
GN DR2473.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
RX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Uterback T., Zalewski C.,
RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RT "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1."
RL Science 286:1571-1577(1999).
CC -!- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.
DR EMBL; AE002076; AAF12016.1; -.
DR PIR; F75270; F75270.
DR TIGR; DR2473; -.
DR GO; GO:0004937; F:monooxygenase activity; IEA.
DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR001128; Cytochrome_P450.
DR Pfam; PF00067; P450; 1.
DR PRINTS; PR00385; P450.
DR PROSITE; PS00086; CYTOCHROME_P450; 1.
KW Heme; Monooxygenase; Oxidoreductase; Complete proteome.
SQ SEQUENCE 381 AA; 41940 MW; F191EA69F1797B53 CRC64;

Query Match 67.7%; Score 44; DB 16; Length 381;
Best Local Similarity 100.0%; Pred. No. 31; Indels 0; Gaps 0;
Matches 9; Conservative 0; Mismatches 0;

QY 1 LPAVVGLSP 9
Db 52 LPAVVGLSP 60

RESULT 3
ID Q8Y280 PRELIMINARY; PRT; 614 AA.
AC Q8Y280;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DE Probable ATP-binding transport ABC transporter protein.
GN RSC0456 OR RS04444.
OS Ralstonia solanacearum (Pseudomonas solanacearum).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Burkholderiaceae; Ralstonia.
OX NCBI_TaxID=305;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=GMT11000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
RA Ariat M., Billault A., Brottier P., Camus J.C., Cattolico L.,
RA Chandler M., Choisme N., Claudel-Renard C., Cunnac S., Demange N.,
RA Gaspin C., Lavie M., Moisan A., Robert C., Saurin W., Schliex T.,
RA Sigquier P., Thebault P., Whalen M., Wincker P., Levy M.,
RA Weissenbach J., Boucher C.A.;
RT "Genome sequence of the plant pathogen Ralstonia solanacearum."
RL Nature 415:497-502(2002).
DR EMBL; AL646059; CAD13984.1; -.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0004009; F:ATP-binding cassette (ABC) transporter acti. .; IEA.

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DR GO; GO:0000166; F:nucleotide binding; IEA.
DR GO; GO:0006810; P:transport; IEA.
DR InterPro; IPR003593; AAA_ATPase.
DR InterPro; IPR003439; ABC_transporter.
DR InterPro; IPR001623; DnaJ_N.
DR Pfam; PF00005; ABC_tran; 1.
DR ProDom; PD000006; ABC_transporter; 1.
DR SMART; SM00382; AAA; 1.
DR PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
DR PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
DR PROSITE; PS00636; DnaJ_1; 1.
KW Complete proteome.
SQ SEQUENCE 614 AA; 69240 MW; E293355B585872142 CRC64;

Query Match 66.9%; Score 43.5; DB 16; Length 614;
Best Local Similarity 45.8%; Pred. No. 63;
Matches 11; Conservative 2; Mismatches 0; Indels 11; Gaps 1;

QY 1 LPAVVG-----LSPGQE 13
Db 501 LPAVLGLLDEVSNWSLSPGQQ 524

RESULT 4
ID Q8HPR8 PRELIMINARY; PRT; 821 AA.
AC Q8HPR8;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE DNA helicase.
OS HEL OR VNG1501G.
ON Halobacterium sp. (strain NRC-1 / ATCC 700922 / JCM 11081).
OC Archaea; Euryarchaeota; Halobacteria; Halobacteriales;
OC Halobacteriaceae; Halobacterium.
OX NCBI_TaxID=64091;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20504483; PubMed=11016950;
RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,
RA Shukla H.D., Lasky S.R., Baliga N.S., Thorsson V., Sbrogna J.,
RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
RA Leithausen B., Keller K., Cruz R., Danson M.J., Hough D.W.,
RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
RA Isenbarger T.A., Peck R.F., Fohlstroder M., Spudich J.L., Jung K.-H.,
RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
RA Ebhardt H., Lowe T.M., Liang P., Riley M., Hood L., Dassarma S.;
RT "Genome sequence of Halobacterium species NRC-1."
RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).
DR EMBL; AE005064; AAG19799.1; -.
DR PIR; C84304; C84304.
DR GO; GO:0004386; F:helicase activity; IEA.
DR GO; GO:0000166; F:nucleotide binding; IEA.
DR GO; GO:0007242; P:intracellular signaling cascade; IEA.
DR InterPro; IPR003593; AAA_ATPase.
DR InterPro; IPR001478; PDZ.
DR SMART; SM00382; AAA; 1.
DR SMART; SM00228; PDZ; 1.
DR KW Helicase; Complete proteome.
SQ SEQUENCE 821 AA; 89848 MW; C454C76B984A5702 CRC64;

Query Match 66.2%; Score 43; DB 17; Length 821;
Best Local Similarity 90.0%; Pred. No. 1e+02;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AVVGLSPGQE 12
Db 326 AVVGLSPAQQ 335

RESULT 5
Q8HXL3 PRELIMINARY; PRT; 1541 AA.
ID Q8HXL3

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AC Q9HXL3;  
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)  
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Hypothetical protein.  
 OS Sus scrofa (Pig).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
 OX NCBI\_TaxID=9823;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=21845885; PubMed=11856879;  
 RA Drogemuller C., Kuiper H., Voss-Nemitz R., Brenig B., Distl O.,  
 RA Leeb T.;  
 RT "Molecular characterization and chromosome assignment of the porcine  
 RT gene COX7A1 coding for the muscle specific cytochrome c oxidase  
 RT subunit VIIa-M";  
 RL Cytogenet. Cell Genet. 94:190-193(2001).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RA Leeb T.;  
 RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ410870; CAD56046.1; -;  
 DR InterPro; IPR001680; WD40.  
 DR Pfam; PF00400; WD40; 12.  
 DR PROSITE; PS50082; WD\_REPEATS\_2; 1.  
 DR PROSITE; PS50294; WD\_REPEATS\_REGION; 3.  
 KW Hypothetical protein.  
 SQ SEQUENCE 1541 AA; 168898 MW; 8188882854AF4F1E CRC64;  
 Query Match 64.6%; Score 42; DB 6; Length 1541;  
 Best Local Similarity 58.3%; Pred. No. 2.9e+02;  
 Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;  
 Qy 1 LPVAVGLSPGEQ 12  
 : : : : :  
 Db 786 MPSEISLSPGEQ 797  
 RESULT 6  
 Q9UXP0  
 AC Q9UXP0 PRELIMINARY; PRT; 326 AA.  
 ID Q9UXP0  
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)  
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE F420-dependent N5,N10-methylene-tetrahydromethanopterin reductase,  
 DE putative.  
 GN FPD.  
 OS Methanobolus tindarius.  
 OC Archaea; Euryarchaeota; Euryarchaeota orders incertae sedis;  
 OC Methanosarcinales; Methanosarcinaceae; Methanobolus.  
 OX NCBI\_TaxID=2221;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=DSM 2278;  
 RA Westenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,  
 RA Gottschalk G., Blaut M.;  
 RT "The F420H2-dehydrogenase from Methanobolus tindarius: Cloning of the  
 RT ffd operon and expression of the genes in Escherichia coli";  
 RL Submitted (SEP-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ011519; CAB56639.1; -;  
 DR PIR; T45226; T45226.  
 DR InterPro; IPR002103; Bac\_luciferase.  
 DR Pfam; PF00296; bac\_luciferase; 1.  
 SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;  
 Query Match 63.1%; Score 41; DB 1; Length 326;  
 Best Local Similarity 70.0%; Pred. No. 88;  
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
 Qy 3 AVVGLSPGEQ 12  
 : : : : :  
 AC Q9HXL3;  
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)  
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Hypothetical protein.  
 OS Sus scrofa (Pig).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
 OX NCBI\_TaxID=9823;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=21845885; PubMed=11856879;  
 RA Drogemuller C., Kuiper H., Voss-Nemitz R., Brenig B., Distl O.,  
 RA Leeb T.;  
 RT "Molecular characterization and chromosome assignment of the porcine  
 RT gene COX7A1 coding for the muscle specific cytochrome c oxidase  
 RT subunit VIIa-M";  
 RL Cytogenet. Cell Genet. 94:190-193(2001).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RA Leeb T.;  
 RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AJ410870; CAD56046.1; -;  
 DR InterPro; IPR001680; WD40.  
 DR Pfam; PF00400; WD40; 12.  
 DR PROSITE; PS50082; WD\_REPEATS\_2; 1.  
 DR PROSITE; PS50294; WD\_REPEATS\_REGION; 3.  
 KW Hypothetical protein.  
 SQ SEQUENCE 1541 AA; 168898 MW; 8188882854AF4F1E CRC64;  
 Query Match 64.6%; Score 42; DB 6; Length 1541;  
 Best Local Similarity 58.3%; Pred. No. 2.9e+02;  
 Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;  
 Qy 1 LPVAVGLSPGEQ 12  
 : : : : :  
 Db 786 MPSEISLSPGEQ 797

Db 88 AILGLGPGEQ 97  
 RESULT 7  
 Q9N158  
 ID Q9N158 PRELIMINARY; PRT; 579 AA.  
 AC Q9N158  
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)  
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Similar to cerebroglycan (Hypothetical protein FLJ38962).  
 GN DKFZP547M109.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 OX NCBI\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Brain;  
 RA Strausberg R.;  
 RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RA Ninomiya K., Wagatsuma M., Kanda K., Kondo H., Yokoi T., Kodaira H.,  
 RA Furuya T., Takahashi M., Kikkawa E., Omura Y., Abe K., Kamihara K.,  
 RA Katsuta N., Sato K., Tanikawa M., Yamazaki M., Sugiyama T., Irie R.,  
 RA Otsuki T., Sato H., Wakamatsu A., Ishii S., Yamamoto J., Isono Y.,  
 RA Kawai-Hiro Y., Saito K., Nishikawa T., Kimura K., Yamashita H.,  
 RA Matsuo K., Nakamura Y., Sekine M., Kikuchi H., Murakawa K.,  
 RA Kanehori K., Takahashi-Fujii A., Oshima A., Sugiyama A., Kawakami B.,  
 RA Suzuki Y., Sugano S., Nagahari K., Masuho Y., Nagai K., Isogai T.;  
 RT "NEDO human cDNA sequencing project";  
 RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Brain;  
 RA Bloecher H., Boecher M., Brandt P., Mewes H.W., Weil B., Wiemann S.;  
 RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; BC027972; AAH27972.1; -;  
 DR EMBL; AK096281; BAC04745.1; -;  
 DR EMBL; AL834418; CAD39080.1; -;  
 DR Genew; HGNC:4450; GPC2.  
 DR GO; GO:0005578; C:extracellular matrix; IEA.  
 DR InterPro; IPR001863; Glypican.  
 DR Pfam; PF01153; Glypican; 1.  
 DR PROSITE; PS01207; GLYPICAN; 1.  
 KW Hypothetical protein.  
 SQ SEQUENCE 579 AA; 62829 MW; 1630E3A22BB83DFA CRC64;  
 Query Match 63.1%; Score 41; DB 4; Length 579;  
 Best Local Similarity 75.0%; Pred. No. 1.6e+02;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 Qy 1 LPVAVGLSPGEQ 12  
 : : : : :  
 Db 439 LPPVVGSPGEQ 450  
 RESULT 8  
 Q9L4W3  
 ID Q9L4W3 PRELIMINARY; PRT; 11096 AA.  
 AC Q9L4W3  
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)  
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE NYSC.  
 GN NYSC.  
 OS Streptomyces noursei.  
 OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
 OC Streptomycinae; Streptomycetaceae; Streptomyces.  
 OX NCBI\_TaxID=1971;  
 RN [1]  
 RP SEQUENCE FROM N.A.

```

RC STRAIN=ATCC 11455;
RX MEDLINE=20334850; PubMed=10873841;
RA Brautaset T., Sekurova O.N., Sletta H., Ellingsen T.E., Strom A.R.,
RA Valla S., Zotchev S.B.;
RT "Biosynthesis of the polyene antifungal antibiotic nystatin in
RT Streptomyces noursei ATCC 11455: analysis of the gene cluster and
RT deduction of the biosynthetic pathway.";
RL Chem. Biol. 7:395-403(2000).
DR EMBL; AF263912; AAF1776.1; -.
DR HSSP; P25715; 1MLA.
DR GO; GO:0004024; F:alcohol dehydrogenase activity, zinc-dependent; IEA.
DR GO; GO:0036740; F:transferase activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR GO; GO:0004314; F:lacyl-carrier protein; IEA.
DR GO; GO:0006633; F:fatty acid biosynthesis; IEA.
DR GO; GO:0008152; P:metabolism; IEA.
DR InterPro; IPR001227; Ac trans.
DR InterPro; IPR002085; Adf_zn_family.
DR InterPro; IPR004410; FdbD.
DR InterPro; IPR000794; Ketoacyl_synth.
DR InterPro; IPR006162; Pantoine S.
DR InterPro; IPR006163; Pp bind.
DR Pfam; PF00698; Acyl_transf; 6.
DR Pfam; PF00107; ADH_Zinc_N; 1.
DR Pfam; PF00109; ketoacyl-synt_C; 6.
DR Pfam; PF02801; ketoacyl-synt_C; 6.
DR Pfam; PF00550; pp-binding; 6.
DR TIGRfams; TIGR00128; fadb; 6.
DR PROSITE; PS00075; ACP DOMAIN; 6.
DR PROSITE; PS00606; B_KETOACYL_SYNTHASE; 6.
DR PROSITE; PS00012; PHOSPHOPANTHEINE; 5.
DR Phosphopantetheine; Transferase.
KW SEQUENCE 11096 AA; 1150415 MW; 776CAEAFCAE551DD CRC64;

Query Match 63.1%; Score 41; DB 2; Length 11096;
Best Local Similarity 63.6%; Pred. No. 3.3e+03;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVGLSPGEQ 12
Db 4998 PEVTGLAPGDQ 5008

RESULT 9
Q8ZYJ3 PRELIMINARY; PRT; 154 AA.
AC Q8ZYJ3;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Hypothetical protein PAE0746.
GN PAE0746.
OS Pyrobaculum aerophilum.
OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;
OC Thermoproteaceae; Pyrobaculum.
OX NCBI_TaxID=13773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IM2 / ATCC 51768 / DSM 7523;
RX MEDLINE=21664397; PubMed=11792869;
RA Fitz-Gibbon S.T., Ladner H., Kim U.-J., Stetter K.O., Simon M.I.,
RA Miller J.H.;
RT "Genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum
RT aerophilum.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989(2002).
DR EMBL; AF009783; AL63000.1; -.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 154 AA; 17403 MW; 1C3D8BCB40324766 CRC64;

Query Match 61.5%; Score 40; DB 17; Length 154;
Best Local Similarity 66.7%; Pred. No. 60;
Matches 10; Conservative 1; Mismatches 2; Indels 2; Gaps 1;

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QY 1 LPAVVGUS--PGEQ 13
Db 32 LPDPVGISYTPGEQ 46

RESULT 10
Q7WCX1 PRELIMINARY; PRT; 242 AA.
AC Q7WCX1;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Bacteriophage-related DNA polymerase.
GN BB3808.
OS Bordetella bronchiseptica (Alcaligenes bronchisepticus).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Alcaligenaceae; Bordetella.
OX NCBI_TaxID=518;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=RB50 / ATCC BAA-588;
RX MEDLINE=22827954; PubMed=12910271;
RA Parkhill J., Sebaihia M., Preston A., Murphy L.D., Thomson N.,
RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,
RA Cerdano-Harraga A.M., Temple L., James K., Harris B., Quail M.A.,
RA Achtman M., Atkin R., Baker S., Basham D., Hason N., Cherevach I.,
RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,
RA Fellwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,
RA Leather S., Moule S., Norberczak H., O'Neill S., Ormond D., Price C.,
RA Rabinowitsch E., Rutter S., Sanders M., Saunders D., Seeger K.,
RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,
RA Unwin I., Whitehead S., Barrell B.G., Maskell D.J.;
RT "Comparative analysis of the genome sequences of Bordetella pertussis,
RT Bordetella parapertussis and Bordetella bronchiseptica.";
RL Nat. Genet. 35:32-40(2003).
DR EMBL; BX640448; CAE35782.1; -.
KW Complete proteome.
SQ SEQUENCE 242 AA; 25627 MW; EECAD9B319823284 CRC64;

Query Match 61.5%; Score 40; DB 16; Length 242;
Best Local Similarity 80.0%; Pred. No. 96;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGEQ 13
Db 96 VVGEAPGEQ 105

RESULT 11
Q8NQA2 PRELIMINARY; PRT; 261 AA.
AC Q8NQA2;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Hypothetical protein Cgl1536.
GN CGL1536.
OS Corynebacterium glutamicum (Brevibacterium flavum).
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Corynebacteriaceae; Corynebacterium.
OX NCBI_TaxID=1718;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA Nakagawa S.;
RT "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP005278; BAB98929.1; -.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 261 AA; 27957 MW; 4D8D51D4DCA3A210 CRC64;

Query Match 61.5%; Score 40; DB 16; Length 261;
Best Local Similarity 53.8%; Pred. No. 1e+02;

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Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 LPVVGLSPGEQ 13  
 Db 137 LPAITVSPGEAD 149

## RESULT 12

Q7W5D6 Q7W5D6 PRELIMINARY; PRT; 298 AA.  
 AC Q7W5D6;  
 DT 01-OCT-2003 (TREMBLrel. 25, Created)  
 DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)  
 DE Bacteriophage-related DNA polymerase.  
 GN BP3357.  
 OS Bordetella parapertussis.  
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;  
 OC Alcaligenaceae; Bordetella.  
 OX NCBI\_TaxID=519;  
 RN [1]

SEQUENCE FROM N.A.  
 RC STRAIN=12822 / ATCC BAA-587;  
 RX MEDLINE=22827954; PubMed=12910271;  
 RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,  
 RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,  
 RA Cerdano-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,  
 RA Achtman M., Atkin R., Baker S., Basham D., Bason N., Cherevach I.,  
 RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,  
 RA Feltwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,  
 RA Leather S., Moule S., Norberczak H., O'Neill S., Ormond D., Price C.,  
 RA Rabinowitsch E., Rutter S., Saunders M., Saunders D., Seeger K.,  
 RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,  
 RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;  
 RT "Comparative analysis of the genome sequences of Bordetella pertussis,  
 RT Bordetella parapertussis and Bordetella bronchiseptica.";  
 RL Nat. Genet. 35:32-40(2003).  
 DR EMBL; EX640433; CAE38642.1; -.  
 KW Complete proteome.  
 SQ SEQUENCE 298 AA; 31202 MW; A858C8199E2AC8B7 CRC64;

Query Match 61.5%; Score 40; DB 16; Length 298;  
 Best Local Similarity 80.0%; Pred. No. 1.2e+02;  
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGEQ 13  
 Db 152 VVGEAPGEQ 161

## RESULT 13

Q7VTI2 Q7VTI2 PRELIMINARY; PRT; 298 AA.  
 AC Q7VTI2;  
 DT 01-OCT-2003 (TREMBLrel. 25, Created)  
 DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)  
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)  
 DE Bacteriophage-related DNA polymerase.  
 GN BP3556.  
 OS Bordetella pertussis.  
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;  
 OC Alcaligenaceae; Bordetella.  
 OX NCBI\_TaxID=520;  
 RN [1]

SEQUENCE FROM N.A.  
 RC STRAIN=Tohama I / ATCC BAA-589 / NCTC 13251;  
 RX MEDLINE=22827954; PubMed=12910271;  
 RA Parkhill J., Sebahia M., Preston A., Murphy L.D., Thomson N.,  
 RA Harris D.E., Holden M.T.G., Churcher C.M., Bentley S.D., Mungall K.L.,  
 RA Cerdano-Tarraga A.M., Temple L., James K., Harris B., Quail M.A.,  
 RA Achtman M., Atkin R., Baker S., Basham D., Bason N., Cherevach I.,  
 RA Chillingworth T., Collins M., Cronin A., Davis P., Doggett J.,  
 RA Feltwell T., Goble A., Hamlin N., Hauser H., Holroyd S., Jagels K.,

RA Leather S., Moule S., Norberczak H., O'Neill S., Ormond D., Price C.,  
 RA Rabinowitsch E., Rutter S., Saunders M., Saunders D., Seeger K.,  
 RA Sharp S., Simmonds M., Skelton J., Squares R., Squares S., Stevens K.,  
 RA Unwin L., Whitehead S., Barrell B.G., Maskell D.J.;  
 RT "Comparative analysis of the genome sequences of Bordetella pertussis,  
 RT Bordetella parapertussis and Bordetella bronchiseptica.";  
 RL Nat. Genet. 35:32-40(2003).  
 DR EMBL; EX640421; CAE43815.1; -.  
 KW Complete proteome.  
 SQ SEQUENCE 298 AA; 31231 MW; 9E4BC84B419506F3 CRC64;

Query Match 61.5%; Score 40; DB 16; Length 298;  
 Best Local Similarity 80.0%; Pred. No. 1.2e+02;  
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 VVGLSPGEQ 13  
 Db 152 VVGEAPGEQ 161

## RESULT 14

Q947A7 Q947A7 PRELIMINARY; PRT; 355 AA.  
 AC Q947A7;  
 DT 01-DEC-2001 (TREMBLrel. 19, Created)  
 DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)  
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)  
 DE Enolase (EC 4.2.1.11) (2-phosphoglycerate dehydratase) (2-phospho-D-  
 DE glycerate hydro-lyase) (Fragment).  
 GN ENO.  
 OS Nitellopsis obtusa.  
 OC Eukaryota; Viridiplantae; Streptophyta; Charales; Characeae;  
 OC Nitellopsis.  
 OX NCBI\_TaxID=40811;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RC STRAIN=X-854;  
 RX MEDLINE=21437986; PubMed=11526220;  
 RA Keeling P.J., Palmer J.D.;  
 RT "Lateral transfer at the gene and subgenic levels in the evolution of  
 RT eukaryotic enolase.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 98:10745-10750(2001).  
 CC -1- CATALYTIC ACTIVITY: 2-PHOSPHO-D-GLYCERATE = PHOSPHOENOLPYRUVATE +  
 CC H(2O).  
 CC -1- COFACTOR: MAGNESIUM IS REQUIRED FOR CATALYSIS AND FOR STABILIZING  
 CC THE DIMER (BY SIMILARITY).  
 CC -1- PATHWAY: GLYCOLYSIS.  
 CC -1- SUBUNIT: HOMODIMER (BY SIMILARITY).  
 CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC (BY SIMILARITY).  
 CC -1- SIMILARITY: BELONGS TO THE ENOLASE FAMILY.  
 DR EMBL; AF348916; AA05455.1; -.  
 DR GO; GO:0000015; C:phosphopyruvate hydratase complex; IEA.  
 DR GO; GO:0016829; F:lyase activity; IEA.  
 DR GO; GO:0000287; F:magnesium ion binding; IEA.  
 DR GO; GO:0004634; F:phosphopyruvate hydratase activity; IEA.  
 DR GO; GO:0006096; P:glycolysis; IEA.  
 DR InterPro; IPR000941; Enolase.  
 DR Pfam; PF00113; enolase; 1.  
 DR Pfam; PF03952; enolase N; 1.  
 DR PRINTS; PR00148; ENOLASE.  
 DR ProDom; PD000902; Enolase; 1.  
 DR PROSITE; PS00164; ENOLASE; 1.  
 DR GlycoLysis; Lyase; Magnesium.  
 FT NON TER 1  
 FT NON TER 355 355  
 SQ SEQUENCE 355 AA; 38295 MW; 400DF160087DE450 CRC64;

Query Match 61.5%; Score 40; DB 10; Length 355;  
 Best Local Similarity 54.5%; Pred. No. 1.4e+02;  
 Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 Db 152 PAVVGLSPGEQ 12



Db 28 PAVIGMDPADQ 38

## RESULT 15

Q8NHX1  
AC Q8NHX1 PRELIMINARY; PRT; 357 AA.  
DT 01-OCT-2002 (Tremblrel. 22, Created)  
DT 01-OCT-2002 (Tremblrel. 22, Last sequence update)  
DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)  
DE Extracellular signal-related kinase lb.  
OS Homo sapiens (Human)  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
SEQUENCE FROM N.A.  
RP Abersold D.M., Yung Y., Seger R.;  
RA "Properties of human ERK1b."  
RT Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.  
DR EMBL: AY033607; RAK52329.1; -;  
DR GO: GO:0005524; F:ATP binding; IEA.  
DR GO: GO:0004707; F:MAP kinase activity; IEA.  
DR GO: GO:0004674; F:protein serine/threonine kinase activity; IEA.  
DR GO: GO:0004713; F:protein-tyrosine kinase activity; IEA.  
DR GO: GO:0016740; F:transferase activity; IEA.  
DR GO: GO:0006468; P:protein amino acid phosphorylation; IEA.  
DR InterPro: IPR008349; Erk\_1\_2\_MAPK.  
DR InterPro: IPR008350; Erk\_3\_4\_MAPK.  
DR InterPro: IPR003527; MAP\_kin.  
DR InterPro: IPR000719; Prot\_kinase.  
DR InterPro: IPR002290; Ser\_thr\_kinase.  
DR InterPro: IPR001245; Tyr\_kinase.  
DR Pfam: PF00069; pkinase; 1.  
DR PRINTS: PR01770; ERKLERK2MAPK.  
DR PRINTS: PR01771; ERK3ERK4MAPK.  
DR ProDom: PD000001; Prot\_kinase; 1.  
DR SMART: SM00220; S\_TKc; 1.  
DR SMART: SM00219; Tyrc; 1.  
DR PROSITE: PS01351; MAPK; 1.  
DR PROSITE: PS00107; PROTEIN\_KINASE\_ATP; 1.  
DR PROSITE: PS00011; PROTEIN\_KINASE\_DOM; 1.  
KW ATP-binding; Kinase; Transferase.  
SQ SEQUENCE 357 AA; 40062 MW; 58C92773983ADA79 CRC64;

Query Match 61.5%; Score 40; DB 4; Length 357;  
Best Local Similarity 72.7%; Pred. No. 1.4e+02;  
Matches 8; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
Db 344 PAAVGLGAGEQ 354

Search completed: May 7, 2004, 12:37:56  
Job time : 25.4767 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 36.27 Seconds

(without alignment)  
101.272 Million cell updates/sec

Title: US-09-786-214A-14

Perfect score: 65

Sequence: 1 LPAVVGSLSPGEQE 13

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

A Genesep29Jan04:\*  
1: Genesep1980s:\*  
2: Genesep1990s:\*  
3: Genesep2000s:\*  
4: Genesep2001s:\*  
5: Genesep2002s:\*  
6: Genesep2003as:\*  
7: Genesep2003bs:\*  
8: Genesep2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	65	100.0	13	3 AAY84268	Aay84268 Peptide d
2	65	100.0	14	3 AAY84266	Aay84266 Peptide d
3	65	100.0	15	3 AAY84269	Aay84269 Peptide d
4	65	100.0	20	3 AAY84265	Aay84265 Truncated
5	65	100.0	25	3 AAY84264	Aay84264 Peptide o
6	61	93.8	13	3 AAY84267	Aay84267 Peptide d
7	45	69.2	234	4 AAB36208	Aab36208 Human imm
8	42	64.6	470	6 AAE34724	Aae34724 Streptomy
9	42	64.6	475	6 AAE34732	Aae34732 Streptomy
10	42	64.6	475	6 AAE34729	Aae34729 Streptomy
11	41	63.1	106	5 ABJ10397	Abj10397 Mutant an
12	41	63.1	106	5 ABJ10395	Abj10395 Mutant an
13	41	63.1	530	7 ADD49105	Add49105 Human NOV
14	41	63.1	549	7 ADD49091	Add49091 Human NOV
15	41	63.1	579	5 ABG70277	Abg70277 Human Gly
16	41	63.1	579	5 ABG97356	Abg97356 Human CGD
17	41	63.1	579	6 ABR39111	Abr39111 Human GPC
18	41	63.1	579	7 ADD49087	Add49087 Human NOV
19	41	63.1	579	7 ADD49107	Add49107 Human NOV
20	41	63.1	579	7 ADD49089	Add49089 Human NOV
21	41	63.1	592	7 ADD49099	Add49099 Human NOV
22	41	63.1	11096	4 AAE10129	Aae10129 Streptomy
23	40	61.5	261	4 AAG91441	Aag91441 C glutami
24	40	61.5	306	4 ABG24698	Abg24698 Novel hum
25	40	61.5	358	4 AAB92530	Aab92530 Human pro

26	40	61.5	359	5 ABB97563	Abb97563 Novel hum
27	40	61.5	428	5 ABB47705	Abb47705 Listeria
28	39	60.0	18	2 AAY41875	Aay41875 Rheumatoi
29	39	60.0	18	4 AAU25388	Aau25388 Schizophr
30	39	60.0	18	4 AAU15732	Aau15732 Schizophr
31	39	60.0	18	5 ABG78871	Abg78871 Multiple
32	39	60.0	23	5 ABP62618	Abp62618 Human imm
33	39	60.0	74	6 ABJ18694	Abj18694 Antibody
34	39	60.0	88	3 AAY56655	Aay56655 Partial p
35	39	60.0	93	5 AAU80983	Aau80983 Human ant
36	39	60.0	94	7 ADD69248	Add69248 Human lig
37	39	60.0	95	6 ABO27154	Abo27154 Human ger
38	39	60.0	95	6 ABO27153	Abo27153 Human ger
39	39	60.0	96	6 ABO27155	Abo27155 Human ger
40	39	60.0	96	6 ABO27150	Abo27150 Human ger
41	39	60.0	100	5 AAE23987	Aae23987 Human MOG
42	39	60.0	104	2 AAW31723	Aaw31723 Alpha lig
43	39	60.0	104	2 AAW26795	Aaw26795 Anti-gp54
44	39	60.0	105	7 ABR61567	Abr61567 HIV-1 neu
45	39	60.0	106	2 AAR41234	Aar41234 Monoclonal

## ALIGNMENTS

RESULT 1  
AAY84268  
ID AAY84268 standard; peptide; 13 AA.  
XX  
AC AAY84268;  
XX  
AC  
DT 12-JUL-2000 (first entry)  
XX  
DE Peptide derived from macrophage colony stimulating gene alternative ORF.  
XX  
KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.  
XX  
XX Synthetic.  
OS Homo sapiens.  
XX  
FN WC200013699-A1.  
XX  
PD 16-MAR-2000.  
XX  
PF 03-SEP-1999; 99WO-US020344.  
XX  
PR 04-SEP-1998; 98US-0099077P.  
XX  
PA (LUDW-) LUDWIG INST CANCER RES.  
XX  
XX  
XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;  
XX WPI; 2000-256859/22.  
XX  
PT Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.  
XX  
XX Example 2; Page 40; 74pp; English.  
XX  
CC The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying

```

CC functional variants and mimetics
XX
SQ Sequence 13 AA;
  Query Match      100.0%; Score 65; DB 3; Length 13;
  Best Local Similarity 100.0%; Pred. No. 0.00057;
  Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGLSPGQE 13
  |||||
Db 1 LPAVVGLSPGQE 13

RESULT 2
AAY84266
ID AAY84266 standard; peptide; 14 AA.
XX
AC AAY84266;
XX
DT 12-JUL-2000 (first entry)
XX
DE Peptide derived from macrophage colony stimulating gene alternative ORF.
XX
KW tumour rejection antigen; macrophage colony stimulating gene;
KW macrophage-colony stimulating factor; antigen presenting cell;
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.
XX
OS Synthetic.
OS Homo sapiens.
XX
FN WO200013699-A1.
XX
PD 16-MAR-2000.
XX
PF 03-SEP-1999; 99WO-US020344.
XX
PR 04-SEP-1998; 98US-0099077P.
XX
PA (LUDW-) LUDWIG INST CANCER RES.
XX
PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;
XX
DR WPI; 2000-256859/22.
XX
PD 16-MAR-2000.
XX
PF 03-SEP-1999; 99WO-US020344.
XX
PR 04-SEP-1998; 98US-0099077P.
XX
PA (LUDW-) LUDWIG INST CANCER RES.
XX
PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;
XX
DR WPI; 2000-256859/22.
XX
PD Isolated polypeptide used to treat subjects having a disorder
PT characterized by expression of alternative open reading frame macrophage-
PT colony stimulating factor comprises 25 amino acid residue sequence.
XX
PS Claim 2; Page 39; 74pp; English.
XX
CC The present sequence represents a peptide which is derived from a tumour
CC rejection antigen precursor encoded by an alternative open reading frame
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from
CC the alternative ORF of macrophage-colony stimulating factor, when
CC presented by an antigen presenting cell having a human leukocyte antigen
CC (HLA) class I molecule, effectively induce the activation and
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic
CC acids derived from the alternate ORF of macrophage-colony stimulating
CC factor are useful for enriching selectively a population of T lymphocytes
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder
CC characterized by expression of the polypeptide, and for identifying
CC functional variants and mimetics
XX
SQ Sequence 14 AA;
  Query Match      100.0%; Score 65; DB 3; Length 14;
  Best Local Similarity 100.0%; Pred. No. 0.00061;
  Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGLSPGQE 13
  |||||
Db 1 LPAVVGLSPGQE 13

RESULT 3
AAY84269
ID AAY84269 standard; peptide; 15 AA.
XX
AC AAY84269;
XX
DT 12-JUL-2000 (first entry)
XX
DE Peptide derived from macrophage colony stimulating gene alternative ORF.
XX
KW tumour rejection antigen; macrophage colony stimulating gene;
KW macrophage-colony stimulating factor; antigen presenting cell;
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.
XX
OS Synthetic.
OS Homo sapiens.
XX
FN WO200013699-A1.
XX
PD 16-MAR-2000.
XX
PF 03-SEP-1999; 99WO-US020344.
XX
PR 04-SEP-1998; 98US-0099077P.
XX
PA (LUDW-) LUDWIG INST CANCER RES.
XX
PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;
XX
DR WPI; 2000-256859/22.
XX
PD Isolated polypeptide used to treat subjects having a disorder
PT characterized by expression of alternative open reading frame macrophage-
PT colony stimulating factor comprises 25 amino acid residue sequence.
XX
PS Example 2; Page 40; 74pp; English.
XX
CC The present sequence represents a peptide which is derived from a tumour
CC rejection antigen precursor encoded by an alternative open reading frame
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from
CC the alternative ORF of macrophage-colony stimulating factor, when
CC presented by an antigen presenting cell having a human leukocyte antigen
CC (HLA) class I molecule, effectively induce the activation and
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic
CC acids derived from the alternate ORF of macrophage-colony stimulating
CC factor are useful for enriching selectively a population of T lymphocytes
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder
CC characterized by expression of the polypeptide, and for identifying
CC functional variants and mimetics
XX
SQ Sequence 15 AA;
  Query Match      100.0%; Score 65; DB 3; Length 15;
  Best Local Similarity 100.0%; Pred. No. 0.00066;
  Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVVGLSPGQE 13
  |||||
Db 3 LPAVVGLSPGQE 15

RESULT 4
AAY84265
ID AAY84265 standard; peptide; 20 AA.
XX
AC AAY84265;
XX
DT 12-JUL-2000 (first entry)
XX
DE Truncated macrophage colony stimulating factor tumour antigen.
KW tumour rejection antigen; macrophage colony stimulating gene;

```

KW macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Homo sapiens.

PN WO200013699-A1.

PD 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

PF 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

PA Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

DR N-PSDB; AAZ99675.

XX Isolated polypeptide used to treat subjects having a disorder

PT characterized by expression of alternative open reading frame macrophage-

PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 3; Page 64; 74pp; English.

XX The present sequence represents a truncated tumour rejection antigen  
 CC precursor, and is encoded by a truncated alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 20 AA;

Query Match 100.0%; Score 65; DB 3; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.0009;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVWGLSPGEQE 13

Db 4 LPAVWGLSPGEQE 16

RESULT 5

AAV84264

ID AAY84264 standard; peptide; 25 AA.

XX AC

XX AAY84264;

DT 12-JUL-2000 (first entry)

XX Peptide of alternate reading frame of macrophage colony stimulating gene.

DE Renal cell carcinoma; antigen; cytotoxic T lymphocyte;

XX tumour rejection antigen; macrophage colony stimulating gene;

KW macrophage-colony stimulating factor; antigen presenting cell;

XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Homo sapiens.

OS WO200013699-A1.

PN 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX

PR 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

DR N-PSDB; AAZ99672.

XX Isolated polypeptide used to treat subjects having a disorder

PT characterized by expression of alternative open reading frame macrophage-

PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 1; Page 64; 74pp; English.

XX The present sequence represents a tumour rejection antigen precursor, and  
 CC is encoded by an alternative open reading frame (ORF) of human macrophage  
 CC colony stimulating gene. Peptides derived from the alternative ORF of  
 CC macrophage-colony stimulating factor, when presented by an antigen  
 CC presenting cell having a human leukocyte antigen (HLA) class I molecule,  
 CC effectively induce the activation and proliferation of CD8+ cytotoxic T  
 CC lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF  
 CC of macrophage-colony stimulating factor are useful for enriching  
 CC selectively a population of T lymphocytes with CD8+ T lymphocytes. They  
 CC are also useful for diagnosing a disorder characterized by expression of  
 CC the polypeptide, and for identifying functional variants and mimetics

XX Sequence 25 AA;

Query Match 100.0%; Score 65; DB 3; Length 25;

Best Local Similarity 100.0%; Pred. No. 0.0012;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LPAVWGLSPGEQE 13

Db 4 LPAVWGLSPGEQE 16

RESULT 6

AAV84267

ID AAY84267 standard; peptide; 13 AA.

XX AC

XX AAY84267;

DT 12-JUL-2000 (first entry)

XX Peptide derived from macrophage colony stimulating gene alternative ORF.

DE tumour rejection antigen; macrophage colony stimulating gene;

XX macrophage-colony stimulating factor; antigen presenting cell;

KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.

OS Homo sapiens.

OS WO200013699-A1.

PN 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder

PT characterized by expression of alternative open reading frame macrophage-

PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 93.8%; Score 61; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0026;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 13  
 |||||  
 Db 1 PAVVGLSPGEQ 12

RESULT 7

AAAB36208  
 ID AAB36208 standard; protein; 234 AA.

AC AAB36208;

DT 15-FEB-2001 (first entry)

XX Human immune system associated protein HISAP-6.

XX Human; immune system associated protein; HISAP-6; immune disorder;  
 KW infection; autoimmune disease; cancer.

XX Homo sapiens.

PN US6135941-A.

XX 24-OCT-2000.

PF 27-MAR-1998; 98US-00049672.

PR 27-MAR-1998; 98US-00049672.

XX (INCY-) INCYTE PHARM INC.

PI Tang YT, Yue H, Lal P, Corley NC, Guegler KJ, Baughn MR;

PI Hillman JL, Au-Young J;

XX WPI: 2001-030926/04.

DR N-PSDB; AAC66524.

XX New human immune system associated proteins (HISAP) and polynucleotides  
 PT encoding the HISAP, useful for diagnosing, treating or preventing immune  
 PT or cell proliferative disorders or infections.

PS Claim 1; Col 59-60; 54pp; English.

XX The present invention provides the coding and protein sequences for a  
 CC number of human immune system associated proteins (HISAPs). These can be  
 CC used in the diagnosis and treatment of various autoimmune disorders,  
 CC infections and cell proliferation diseases. The diseases include AIDS,  
 CC adult respiratory distress syndrome, anaemia, asthma, atherosclerosis,  
 CC Crohn's disease, irritable bowel syndrome, multiple sclerosis, myasthenia  
 CC gravis, osteoarthritis, rheumatoid arthritis, scleroderma, systemic lupus  
 CC erythematosus, arteriosclerosis, cirrhosis and cancer

XX Sequence 234 AA;

Query Match 69.2%; Score 45; DB 4; Length 234;  
 Best Local Similarity 72.7%; Pred. No. 26;  
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 |||||  
 Db 28 PAIVSLSPGER 38

RESULT 8

AAE34724  
 ID AAE34724 standard; protein; 470 AA.

XX AAE34724;

DT 14-MAY-2003 (first entry)

XX Streptomyces rimosus ema3 protein.

XX P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
 KW emamectin; insecticide; ema3 protein.

XX Streptomyces rimosus.

PN WO200292801-A2.

PD 21-NOV-2002.

PF 15-MAY-2002; 2002WO-EP005363.

PR 16-MAY-2001; 2001US-0291149P.

XX (SYGN) SYNGENTA PARTICIPATIONS AG.

PI Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;

DR WPI: 2003-140280/13.

DR N-PSDB; AAD53019.

XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.

PS Claim 17; Page 107-108; 157pp; English.

XX The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus ema3 protein

XX Sequence 470 AA;

Query Match 64.6%; Score 42; DB 6; Length 470;  
 Best Local Similarity 72.7%; Pred. No. 1.8e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 LPAVVGLSPGE 11  
 |||||  
 Db 44 LPSYVGLHPGE 54

RESULT 9

AAE34732  
 ID AAE34732 standard; protein; 475 AA.

XX AAE34732;

DT 14-MAY-2003 (first entry)  
 XX Streptomyces rimosus emal1 protein.  
 DE P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
 KW emamectin; insecticide; emal1 protein.  
 XX Streptomyces rimosus.  
 OS WO200292801-A2.  
 XX 21-NOV-2002.  
 XX 15-MAY-2002; 2002WO-EP005363.  
 XX 16-MAY-2001; 2001US-0291149P.  
 XX (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 PA Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; AAD53027.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 121-122; 157pp; English.  
 CC The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus emal1 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 64.6%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. NO. 1.8e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 DB 49 LPSYVGLHPGE 59  
 ||: ||| |||  
 RESULT 10  
 AAEE34729  
 ID AAEE34729 standard; protein; 475 AA.  
 AC AAEE34729;  
 XX 14-MAY-2003 (first entry)  
 DE Streptomyces albofaciens emas8 protein.  
 KW P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
 KW emamectin; insecticide; emas8 protein.  
 XX Streptomyces albofaciens.  
 OS WO200292801-A2.  
 XX 21-NOV-2002.  
 XX 15-MAY-2002; 2002WO-EP005363.  
 XX 16-MAY-2001; 2001US-0291149P.  
 XX (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 PA Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; AAD53027.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 121-122; 157pp; English.  
 CC The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus emal1 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 64.6%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. NO. 1.8e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 DB 49 LPSYVGLHPGE 59  
 ||: ||| |||  
 RESULT 10  
 AAEE34729  
 ID AAEE34729 standard; protein; 475 AA.  
 AC AAEE34729;  
 XX 14-MAY-2003 (first entry)  
 DE Streptomyces albofaciens emas8 protein.  
 KW P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
 KW emamectin; insecticide; emas8 protein.  
 XX Streptomyces albofaciens.  
 OS WO200292801-A2.  
 XX 21-NOV-2002.  
 XX 15-MAY-2002; 2002WO-EP005363.  
 XX 16-MAY-2001; 2001US-0291149P.  
 XX (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 PA Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; AAD53024.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 116-117; 157pp; English.  
 CC The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces albofaciens emas8 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 64.6%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. NO. 1.8e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 DB 49 LPSYVGLHPGE 59  
 ||: ||| |||  
 RESULT 11  
 ABJ10397  
 ID ABJ10397 standard; protein; 106 AA.  
 AC ABJ10397;  
 XX 28-NOV-2002 (first entry)  
 DE Mutant anti-mesothelin Fv (ST6) variable light chain.  
 KW Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.  
 OS Unidentified.  
 OS Synthetic.  
 XX WO200240545-A2.  
 XX 23-MAY-2002.  
 XX 16-NOV-2001; 2001WO-US043602.  
 XX 17-NOV-2000; 2000US-0249805P.  
 XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.  
 XX Pastan IH, Onda M, Nagata S, Tsutsumi Y, Vincent JJ, Kreitman RJ;  
 PI Vasmatzis G, Lee B;  
 XX WPI; 2002-500208/53.  
 DR N-PSDB; AET08085.  
 XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.  
 XX Claim 12; Fig 2; 82pp; English.

XX PA (SYGN ) SYNGENTA PARTICIPATIONS AG.  
 XX Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 XX Buckel TG;  
 XX WPI; 2003-140280/13.  
 DR N-PSDB; AAD53024.  
 XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.  
 XX Claim 17; Page 116-117; 157pp; English.  
 CC The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces albofaciens emas8 protein  
 XX Sequence 475 AA;  
 SQ  
 Query Match 64.6%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. NO. 1.8e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 LPAVVGLSPGE 11  
 DB 49 LPSYVGLHPGE 59  
 ||: ||| |||  
 RESULT 11  
 ABJ10397  
 ID ABJ10397 standard; protein; 106 AA.  
 AC ABJ10397;  
 XX 28-NOV-2002 (first entry)  
 DE Mutant anti-mesothelin Fv (ST6) variable light chain.  
 KW Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.  
 OS Unidentified.  
 OS Synthetic.  
 XX WO200240545-A2.  
 XX 23-MAY-2002.  
 XX 16-NOV-2001; 2001WO-US043602.  
 XX 17-NOV-2000; 2000US-0249805P.  
 XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.  
 XX Pastan IH, Onda M, Nagata S, Tsutsumi Y, Vincent JJ, Kreitman RJ;  
 PI Vasmatzis G, Lee B;  
 XX WPI; 2002-500208/53.  
 DR N-PSDB; AET08085.  
 XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.  
 XX Claim 12; Fig 2; 82pp; English.

CC The invention comprises the amino acid and coding sequences of  
 CC recombinant immunotoxin proteins. The immunotoxin proteins of the  
 CC invention contain an antibody (or an antigen-binding fragment) with a  
 CC substitution of a negatively charged amino acid for an uncharged or  
 CC positively charged amino acid. The immunotoxins of the invention have  
 CC reduced liver toxicity. The immunotoxins of the invention are useful for  
 CC killing a malignant cell (e.g. a cancer cell). The present amino acid  
 CC sequence represents a recombinant immunotoxin of the invention  
 XX  
 SQ Sequence 106 AA;

Query Match 63.1%; Score 41; DB 5; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 51;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 DB 8 PAIMASPGEQ 18  
 ||: |||||

## RESULT 12

ABJ10395  
 ID ABJ10395 standard; protein; 106 AA.

AC ABJ10395;

XX  
 XT 28-NOV-2002 (first entry)

DE Mutant anti-Tac Fv (M16) variable light chain.

XX Mutant; mutein; recombinant immunotoxin; reduced liver toxicity; cancer.

XX Unidentified.

OS Synthetic.

XX WO200240545-A2.

XX 23-MAY-2002.

XX 16-NOV-2001; 2001WO-US043602.

XX 17-NOV-2000; 2000US-0249805P.

XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.

XX Pastan IH, Onda M, Nagata S, Tsutsumi Y, Vincent JJ, Kreitman RJ;  
 PI Vasmatazis G, Lee B;

XX WPI; 2002-500208/53.

DR N-PSDB; ABT08083.

XX Recombinant immunotoxins with reduced liver toxicity for killing  
 PT malignant cells bearing antigen comprising mutated framework regions of  
 PT antibody heavy and/or light chain or antigen-binding portion of  
 PT immunotoxin.

XX Claim 8; Fig 2; 82pp; English.

XX The invention comprises the amino acid and coding sequences of  
 CC recombinant immunotoxin proteins. The immunotoxin proteins of the  
 CC invention contain an antibody (or an antigen-binding fragment) with a  
 CC substitution of a negatively charged amino acid for an uncharged or  
 CC positively charged amino acid. The immunotoxins of the invention have  
 CC reduced liver toxicity. The immunotoxins of the invention are useful for  
 CC killing a malignant cell (e.g. a cancer cell). The present amino acid  
 CC sequence represents a recombinant immunotoxin of the invention  
 XX  
 SQ Sequence 106 AA;

Query Match 63.1%; Score 41; DB 5; Length 106;  
 Best Local Similarity 63.6%; Pred. No. 51;  
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 PAVVGLSPGEQ 12  
 DB 8 PAIMASPGEQ 18  
 ||: |||||

## RESULT 13

ADD49105  
 ID ADD49105 standard; protein; 530 AA.

XX AC ADD49105;

XX 15-JAN-2004 (first entry)

XX Human NOV15k SEQ ID 78.

XX Antidiabetic; anorectic; cardiac; hypotensive; antiarteriosclerotic;  
 KW viricide; antibacterial; fungicide; protozoacide; nootropic;  
 KW neuroprotective; antiparkinsonian; anticonvulsant; osteopathic;  
 KW antiarthritic; antiinflammatory; dermatological; antiasthmatic;  
 KW antilipemic; gene therapy; NOV protein; metabolic disorder; diabetes;  
 KW obesity; viral infection; bacterial infection; fungal infection;  
 KW helminthic infection; protozoal infection; anorexia; cancer;  
 KW cardiovascular disease; hypertension; atherosclerosis;  
 KW neurodegenerative disorder; Alzheimer's disease; Parkinson's disease;  
 KW epilepsy; immune disorder; osteoarthritis; haematopoietic disorder;  
 KW inflammatory skin disorder; asthma; dyslipidemia; human.

OS Homo sapiens.

XX WO2003060149-A2.

XX 24-JUL-2003.

XX 06-JAN-2003; 2003WO-US000252.

XX 04-JAN-2002; 2002US-0345222P.

XX 14-JAN-2002; 2002US-0348693P.

XX 16-JAN-2002; 2002US-0349182P.

XX 17-JAN-2002; 2002US-0349733P.

XX 18-JAN-2002; 2002US-0350263P.

XX 24-JAN-2002; 2002US-0351977P.

XX 28-MAY-2002; 2002US-0383758P.

XX 05-JUN-2002; 2002US-0385969P.

XX 11-JUN-2002; 2002US-0387834P.

XX 17-JUL-2002; 2002US-0396407P.

XX 30-SEP-2002; 2002US-0415115P.

XX 03-JAN-2003; 2003US-00336603.

XX (CURA-) CURAGEN CORP.

XX Grose WM, Alsobrook JP, Anderson DW, Burgess CE, Edinger SR;  
 PI Ellerman K, Furtak K, Gangolli EA, Gerlach VL, Gilbert JA;

PI Gunther E, Gorman L, Guo X, Ji W, Li L, Miller CE, Padigar M;  
 PI Patturajan M, Rastelli L, Macdougall JR, Mishra VS, Smithson G;

PI Spytek KA, Stone DJ, Shenoy SG, Taupier RJ, Vernet CAM, Zhong M;  
 PI Malyankar UM, Millet I, Kekuda R;

XX WPI; 2003-587288/55.

DR N-PSDB; ADD49104.

XX New isolated NOVX polypeptides and polynucleotides, useful for  
 PT preventing, diagnosing or treating NOVX-associated disorders, e.g.  
 PT osteoarthritis, obesity, atherosclerosis, cancer, Parkinson's disease,  
 PT asthma, or infections.

XX Claim 1; Page 181-182; 31pp; English.

XX The present invention relates to novel NOV proteins and their coding  
 CC sequences (ADD49028-ADD49131). The proteins and coding sequences are  
 CC useful in the manufacture of a medicament for treating a syndrome  
 CC associated with a human disease, preferably a NOV-associated disorder  
 CC such as metabolic disorders, diabetes, obesity, infectious diseases  
 CC (viral, bacterial, fungal, helminthic, and protozoal), anorexia, cancer,

CC cardiovascular diseases (hypertension, atherosclerosis),  
 CC neurodegenerative disorders, Alzheimer's disease, Parkinson's disease,  
 CC epilepsy, immune disorders (osteoarthritis), hematopoietic disorders,  
 CC inflammatory skin disorders, asthma and various dyslipidemias. The coding  
 CC sequences and proteins may also be used as targets for the identification  
 CC of small molecules that modulate or inhibit e.g. neurogenesis, cell  
 CC differentiation, cell proliferation, haematopoiesis, wound healing and  
 CC angiogenesis, in gene therapy, in generation of antibodies that bind  
 CC immunospecifically to NOV substances for use in therapeutic or diagnostic  
 CC methods.  
 XX  
 XX Sequence 530 AA;

Query Match 63.1%; Score 41; DB 7; Length 530;

Best Local Similarity 75.0%; Pred. No. 2.9e+02;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 LPAVVGSLSPGEQ 12

Db 416 LPPVVGSPAEQ 427

RESULT 14

ADD49091

ID ADD49091 standard; protein; 549 AA.

XX ADD49091;

XX 15-JAN-2004 (first entry)

XX Human NOV15d SEQ ID 64.

XX Antidiabetic; anorectic; cardiatic; hypotensive; antiarteriosclerotic;  
 XX virucide; antibacterial; fungicide; protozoacide; nootropic;  
 XX neuroprotective; antiparkinsonian; anticonvulsant; osteopathic;  
 XX antiarthritic; antiinflammatory; dermatological; antiasthmatic;  
 XX antilipemic; gene therapy; NOV protein; metabolic disorder; diabetes;  
 XX obesity; viral infection; bacterial infection; fungal infection;  
 XX helminthic infection; protozoal infection; anorexia; cancer;  
 XX cardiovascular disease; hypertension; atherosclerosis;  
 XX neurodegenerative disorder; Alzheimer's disease; Parkinson's disease;  
 XX epilepsy; immune disorder; osteoarthritis; haematopoietic disorder;  
 XX inflammatory skin disorder; asthma; dyslipidemia; human.

XX Homo sapiens.

XX WO2003060149-A2.

XX 24-JUL-2003.

XX 06-JAN-2003; 2003WO-US000252.

XX 04-JAN-2002; 2002US-0345222P.

XX 14-JAN-2002; 2002US-0348693P.

XX 16-JAN-2002; 2002US-0349182P.

XX 17-JAN-2002; 2002US-0349733P.

XX 18-JAN-2002; 2002US-0350263P.

XX 24-JAN-2002; 2002US-0351977P.

XX 28-MAY-2002; 2002US-0383758P.

XX 05-JUN-2002; 2002US-0385969P.

XX 11-JUN-2002; 2002US-0387834P.

XX 17-JUL-2002; 2002US-0396407P.

XX 30-SEP-2002; 2002US-0415115P.

XX 03-JAN-2003; 2003US-00336603.

XX (CURA-) CURAGEN CORP.

XX Groesse WM, Alsobrook JP, Anderson DW, Burgess CE, Edinger SR;

XX Ellerman K, Furtak K, Gangolli EA, Gerlach VL, Gilbert JA;

XX Gunther E, Gorman L, Guo X, Ji W, Li L, Miller CE, Padigaru M;

XX Patturajan M, Rastelli L, Macdougall JR, Mishra VS, Smithson G;

XX Spytek KA, Stone DJ, Shenoy SG, Taupier RJ, Vernet CM, Zhong M;

XX Malyankar UM, Millet I, Kekuda R;

XX WPI; 2003-587288/55.  
 XX N-PSDB; ADD49090.  
 XX New isolated NOVX polypeptides and polynucleotides, useful for  
 PT preventing, diagnosing or treating NOVX-associated disorders, e.g.  
 PT osteoarthritis, obesity, atherosclerosis, cancer, Parkinson's disease,  
 PT asthma, or infections.  
 XX  
 PS Claim 1; Page 177; 311pp; English.

XX The present invention relates to novel NOV proteins and their coding  
 CC sequences (ADD49028-ADD49131). The proteins and coding sequences are  
 CC useful in the manufacture of a medicament for treating a syndrome  
 CC associated with a human disease, preferably a NOV-associated disorder  
 CC such as metabolic disorders, diabetes, obesity, infectious diseases  
 CC (viral, bacterial, fungal, helminthic, and protozoal), anorexia, cancer,  
 CC cardiovascular diseases (hypertension, atherosclerosis),  
 CC neurodegenerative disorders, Alzheimer's disease, Parkinson's disease,  
 CC epilepsy, immune disorders (osteoarthritis), hematopoietic disorders,  
 CC inflammatory skin disorders, asthma and various dyslipidemias. The coding  
 CC sequences and proteins may also be used as targets for the identification  
 CC of small molecules that modulate or inhibit e.g. neurogenesis, cell  
 CC differentiation, cell proliferation, haematopoiesis, wound healing and  
 CC angiogenesis, in gene therapy, in generation of antibodies that bind  
 CC immunospecifically to NOV substances for use in therapeutic or diagnostic  
 CC methods.

XX Sequence 549 AA;

Query Match 63.1%; Score 41; DB 7; Length 549;

Best Local Similarity 75.0%; Pred. No. 3e+02;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 LPAVVGSLSPGEQ 12

Db 419 LPPVVGSPAEQ 430

RESULT 15

ABG70277

ID ABG70277 standard; protein; 579 AA.

XX ABG70277;

XX 05-NOV-2002 (first entry)

XX Human Glypican-2 Precursor-like protein #1.

XX Human; NOVX; pathological condition; NOVX-associated disorder;  
 XX von Hippel-Lindau syndrome; cirrhosis; transplantation disorder;  
 XX pancreatitis; obesity; diabetes; autoimmune disease; infertility;  
 XX renal artery stenosis; interstitial nephritis; glomerulonephritis;  
 XX polycystic kidney disease; cataract; Alzheimer's disease; cancer;  
 XX acoustic trauma; cardiomyopathy; atherosclerosis; hypertension;  
 XX congenital heart defect; scleroderma; endometriosis; haemophilia;  
 XX dementia; stroke; Parkinson's disease; Huntington's disease; epilepsy;  
 XX multiple sclerosis; anxiety; pain; leukaemia; hypothyroidism; psoriasis;  
 XX acne; wound; asthma; human disease; calpain; epsin; zinc finger;  
 XX low density lipoprotein B; LDLB; purinoceptor; CG8841; synaptotagmin;  
 XX serine protease IL8P; mitogen activated protein kinase kinase-2;  
 XX glypican-2 precursor; thymosin beta-10.

XX Homo sapiens.

XX WO200255702-A2.

XX 18-JUL-2002.

XX 26-OCT-2001; 2001WO-US050925.

XX 26-OCT-2000; 2000US-0243320P.

XX 26-OCT-2000; 2000US-0243592P.



PR 26-OCT-2000; 2000US-0243642P.  
 PR 27-OCT-2000; 2000US-0243681P.  
 PR 27-OCT-2000; 2000US-0243681P.  
 PR 31-OCT-2000; 2000US-0244443P.  
 PR 01-NOV-2000; 2000US-0244995P.  
 PR 01-NOV-2000; 2000US-0245029P.  
 PR 02-NOV-2000; 2000US-0245293P.  
 PR 02-NOV-2000; 2000US-0245315P.  
 PR 02-NOV-2000; 2000US-0245316P.  
 PR 19-JAN-2001; 2001US-0262994P.  
 PR 15-FEB-2001; 2001US-0269056P.  
 PR 02-MAR-2001; 2001US-0272923P.  
 PR 15-MAR-2001; 2001US-0276565P.  
 PR 07-SEP-2001; 2001US-0318119P.  
 XX (CURA-) CURAGEN CORP.  
 PA  
 XX Gangolli EA, Spytek KA, Gilbert J, Casman S, Blalock A, Li L;  
 PI Vernet CAM, Shency S, Mishra V, Furtak K, Gerlach V, Edinger S;  
 PI Malyankar U, Stone D, Millet I, Smithson G, Gunther E, Padigaru M;  
 PI Taupier RJ, Anderson D;  
 XX  
 DR WPI; 2002-590673/63.  
 DR N-PSDB; ABK51684.  
 XX  
 PT Isolated NOVX polypeptides and nucleic acid molecules useful for  
 PT treating, preventing, diagnosing and researching pathological conditions  
 PT in humans with a NOVX-associated disorders, e.g. cancer, stroke or  
 PT Alzheimer's disease.  
 XX  
 PS Claim 1; Page 57; 236pp; English.  
 XX  
 CC The present invention relates to a new polypeptide that comprises any of  
 CC 17 fully defined sequences of 43-990 amino acids given in the  
 CC specification. The NOVX polypeptide, nucleic acid and antibody of the  
 CC invention are useful for treating or preventing a pathological condition  
 CC in humans with a NOVX-associated disorder, e.g. Von Hippel-Lindau  
 CC syndrome, cirrhosis, transplantation disorders, pancreatitis, obesity,  
 CC diabetes, autoimmune disease, renal artery stenosis, interstitial  
 CC nephritis, glomerulonephritis, polycystic kidney disease, cataract,  
 CC Alzheimer's disease, acoustic trauma, cancer, infertility, heart  
 CC cardiomyopathies, atherosclerosis, hypertension, congenital heart  
 CC defects, scleroderma, endometriosis, haemophilia, dementia, stroke,  
 CC Parkinson's disease, Huntington's disease, epilepsy, multiple sclerosis,  
 CC anxiety, pain, leukaemia, hypothyroidism, psoriasis, acne, wounds and  
 CC asthma. They are also useful for the manufacture of a medicament for  
 CC treating a syndrome associated with a human disease, specifically a NOVX-  
 CC associated disorder. They may also be useful in therapeutic applications  
 CC including protein therapy, as small molecule drug targets, as antibody  
 CC targets, as diagnostic and/or prognostic markers, in gene therapy, as  
 CC research tools and in tissue regeneration. The present amino acid  
 CC sequence represents one of the 17 novel proteins of the invention  
 XX  
 SQ Sequence 579 AA;  
 Query Match 63.1%; Score 41; DB 5; Length 579;  
 Best Local Similarity 75.0%; Pred. No. 3.2e+02;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 Qy 1 LPVAVGLSPGEQ 12  
 Db 439 LPPVVGGSPEAQ 450  
 Search completed: May 7, 2004, 12:33:46  
 Job time : 36.27 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:20:30 ; Search time 5.4 Seconds  
(without alignments)  
144.639 Million cell updates/sec

Title: US-09-786-214A-15

Perfect score: 75

Sequence: 1 AGLPAVGLSPGEQE 15

Scoring table: BLOSUM62

Gapop 10.0 ; Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_42:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	57.3	658	1	VG18_BPT4
2	42	56.0	326	1	MER_METTI
3	42	56.0	508	1	COBI_MYCTU
4	41	54.7	156	1	RUVX_CAUCR
5	41	54.7	213	1	MDCG_XANAC
6	41	54.7	520	1	AT15_YEAST
7	41	54.7	774	1	LOL2_HUMAN
8	40	53.3	157	1	RISB_PYRPU
9	40	53.3	359	1	ALF2_PEA
10	40	53.3	359	1	ALF2_PEA
11	40	53.3	361	1	VAL1_TMOV
12	40	53.3	633	1	PLB5_SCHPO
13	40	53.3	773	1	YHGF_ECOLI
14	40	53.3	813	1	CADM_MOUSE
15	40	53.3	2364	1	PCSA_BOVIN
16	39	52.0	115	1	KV31_HUMAN
17	39	52.0	115	1	KV51_MOUSE
18	39	52.0	146	1	DTD_XANAC
19	39	52.0	385	1	Y53_METUA
20	39	52.0	597	1	NR41_RAT
21	39	52.0	1402	1	N160_MOUSE
22	39	52.0	1596	1	GLI3_HUMAN
23	39	52.0	1636	1	RUD3_YEAST
24	38	50.7	146	1	DTD_XANCP
25	38	50.7	299	1	DAF1_STRCO
26	38	50.7	299	1	NUCG_BOVIN
27	38	50.7	331	1	ILVC_ANASP
28	38	50.7	344	1	ILVC_OCEIH
29	38	50.7	390	1	COBL_MYCTU
30	38	50.7	395	1	HNEB_ARCFU
31	38	50.7	429	1	R31_LEULA
32	38	50.7	484	1	COBQ_PSEDE
33	38	50.7	507	1	CATA_PICAN

RESULT 1  
VG18\_BPT4  
ID VG18\_BPT4 STANDARD; PRT; 658 AA.  
AC P13332; Q9T0U3;  
DT 01-JAN-1990 (Rel. 13, Created)  
DT 28-FEB-2003 (Rel. 41, Last sequence update)  
DT 10-OCT-2003 (Rel. 42, Last annotation update)  
DE Tail sheath protein Gp18.  
GN 18.  
OS Bacteriophage T4.  
OC Viruses; dsDNA viruses, no RNA stage; Caudovirales; Myoviridae;  
OC T4-like viruses.  
OX NCBI\_TaxID=10665;  
RN [1]  
SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.  
RP MEDLINE=8815753; PubMed=2964531;  
RA Arisaka F., Nakako T., Takahashi H., Ishii S.-I.;  
RT "Nucleotide sequence of the tail sheath gene of bacteriophage T4 and  
RT amino acid sequence of its product.";  
RL J. Virol. 62:1186-1193(1988).  
RN [2]  
SEQUENCE FROM N.A.  
RP MEDLINE=22514363; PubMed=12626685;  
RA Miller E.S., Kutter E., Mosig G., Arisaka F., Kunisawa T., Ruger W.;  
RT "Bacteriophage T4 genome.";  
RL Microbiol. Mol. Biol. Rev. 67:86-156(2003).  
RN [3]  
SEQUENCE OF 638-658 FROM N.A.  
RA Arisaka F., Ishimoto L., Kasaavetis G., Kumazaki T., Ishii S.-I.;  
RT "Nucleotide sequence of the tail tube structural gene of  
RT bacteriophage T4.";  
RL J. Virol. 62:882-886(1988).  
CC -!- FUNCTION: The contractile tail of bacteriophage T4 consists of a  
CC contractile sheath, a tube and a baseplate. 144 protomers of Gp18,  
CC arranged in 24 annuli, form the contractile tail sheath.  
CC -!- SUBUNIT: Monomer.  
CC -!- SIMILARITY: 30% IDENTITY TO THE TAIL SHEATH PROTEIN OF PHAGE P817,  
CC AND BY EXTENSION, TO P.AERUGINOSA R-TYPE PYOCINS (ANTIBIOTICS).  
CC -----  
CC THIS SWISS-PROT entry is copyright. It is produced through a collaboration  
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CC use by non-profit institutions as long as its content is in no way  
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CC or send an email to license@isb-sib.ch).  
CC -----  
DR EMBL; M19085; AAA32541.1; -;  
DR EMBL; AF158101; AAD42423.1; -;  
DR PIR; JF0021; GKBP74.  
DR InterPro; IPR007067; Phage sheath 1.  
DR Pfam; PF04984; Phage\_sheath\_1; 1.  
DR Structural protein.  
FT INIT MET 0 175 SER-RICH.  
FT DOMAIN 171 175

Q9v4t3 drosophila  
P22361 mus musculus  
P15257 rattus norv  
P20823 homo sapien  
P43925 haemophilus  
P57938 pasteurella  
Q9krb2 vibrio chol  
O97952 macaca fasc  
O97960 papio hamad  
P19091 mus musculus  
P15207 rattus norv  
Q9tt90 canis faml

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FT CONFLICT 99 99 D -> E (IN REF. 1).
FT CONFLICT 147 150 GKNY -> AKII (IN REF. 1).
FT CONFLICT 300 300 E -> G (IN REF. 1).
FT CONFLICT 398 398 A -> V (IN REF. 1).
FT CONFLICT 453 453 H -> Y (IN REF. 1).
FT CONFLICT 594 594 N -> I (IN REF. 1).
SQ SEQUENCE 658 AA; 71199 MW; 6E045F40D39AF21D CRC64;

Query Match
Best Local Similarity 57.3%; Score 43; DB 1; Length 658;
Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPGE 13
DB 217 GIFGVVALYFGE 228

RESULT 2
MER_METTII
ID MER_METTII STANDARD; PRT; 326 AA.
AC Q9UXP0;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Coenzyme F420-dependent N(5),N(10)-methylentetrahydromethanopterin
DE reductase (EC 1.5.99.11) (Methylene-H(4)MPT reductase).
GN MER OR PFDA.
OS Methanobolus tindarius.
OC Archaea; Euryarchaeota; Methanomicrobia; Methanosarcinales;
OC Methanosarcinaceae; Methanobolus.
OX NCBI_TaxID=2221;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM 2278;
RX MEDLINE=99132696; PubMed=9933933;
RA Westenberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,
RA Gottschalk G., Blaut M.;
RA "The F420H2-dehydrogenase from Methanobolus tindarius: cloning of the
RA ffd operon and expression of the genes in Escherichia coli.";
RA FEMS Microbiol. Lett. 170:389-398(1999).
CC -!- FUNCTION: Catalyzes the reversible reduction of methylene-H(4)MPT
CC to methyl-H(4)MPT (By similarity).
CC -!- CATALYTIC ACTIVITY: N(5),N(10)-methylentetrahydromethanopterin +
CC reduced coenzyme F420 = 5-methyl-5,6,7,8-tetrahydromethanopterin +
CC coenzyme F420.
CC -!- PATHWAY: Methanogenesis from carbon dioxide; fifth step.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the mer family.

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or send an email to license@isb-sib.ch).

EMBL; AJ011519; CAB56639.1; -.
DR PIR; T45226; T45226.
DR HAMAP; MF 01091; -.
DR InterPro; IPR002103; Bac_luciferase.
DR Pfam; PF00296; Bac_luciferase; 1.
KW Methanogenesis; One-carbon metabolism; Oxidoreductase.
SQ SEQUENCE 326 AA; 34043 NW; 16F3AB9733A45D82 CRC64;

Query Match
Best Local Similarity 56.0%; Score 42; DB 1; Length 326;
Matches 8; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGLPAVVGLSPGEQ 14
DB 84 SGGRAILGLGFGEQ 97

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RESULT 3
COBI_MYCTTU
ID COBI_MYCTTU STANDARD; PRT; 508 AA.
AC Q10677;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cobalamin biosynthesis protein COBIJ [Includes: Precorrin-2 C20-
DE methyltransferase (EC 2.1.1.130) (S-adenosyl-L-methionine-precorrin-2
DE methyltransferase) (SP2MT); Precorrin-3 methylase (EC 2.1.1.-)].
GN COBIJ OR COBI OR RV2066 OR MT2126 OR MTCY49.05 OR MB2092.
OS Mycobacterium tuberculosis, and
OS Mycobacterium bovis.
OC Bacteria; Actinobacteria; Actinomycetidae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773, 1765;
RN [1]
RP SEQUENCE FROM N.A.
RC SPECIES=M.tuberculosis; STRAIN=H37Rv;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RA complete genome sequence.";
RA Nature 393:537-544(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC SPECIES=M.tuberculosis; STRAIN=CDC 1551 / Oshkosh;
RX MEDLINE=22206494; PubMed=12218036;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W., Jacobs W.R. Jr., Venter J.C., Fraser C.M.;
RA "Whole-genome comparison of Mycobacterium tuberculosis clinical and
RA laboratory strains.";
RA J. Bacteriol. 184:5479-5490(2002).
RN [3]
RP SEQUENCE FROM N.A.
RC SPECIES=M.bovis; STRAIN=AF2122/97;
RX MEDLINE=22709107; PubMed=12788972;
RA Garnier T., Eigmeier K., Camus J.-C., Medina N., Mansoor H.,
RA Pryor M., Duthoy S., Grondin S., Lacroix C., Monsemp C., Simon S.,
RA Harris B., Atkin R., Doggett J., Mayes R., Keating L., Wheeler P.R.,
RA Parkhill J., Barrell B.G., Cole S.T., Gordon S.V., Hewinson R.G.;
RA "The complete genome sequence of Mycobacterium bovis.";
RA Proc. Natl. Acad. Sci. U.S.A. 100:7877-7882(2003).
CC -!- FUNCTION: METHYLATES PRECORRIN-2 AT THE C-20 POSITION TO PRODUCE
CC PRECORRIN-3A (BY SIMILARITY).
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + precorrin-2 = S-
CC adenosyl-L-homocysteine + precorrin-3A.
CC -!- PATHWAY: Cobalamin biosynthesis.
CC -!- SIMILARITY: BELONGS TO A FAMILY THAT GROUPS SUMT, CYSG, CBIF/COBM
CC AND CBIL/COBI.

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or send an email to license@isb-sib.ch).

EMBL; Z73966; CAA98214.1; -.
DR EMBL; AE007063; AAK46406.1; -.
DR EMBL; BX248341; CAD96945.1; -.

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DR PIR; E70764; E70764.
DR TIGR; MT2126; -.
DR TubercuList; RV2066; -.
DR InterPro; IPR006364; Cobi ChlL.
DR InterPro; IPR006363; CobiJ.
DR InterPro; IPR008878; Cox/pox Metransf.
DR InterPro; IPR003043; Uropor_Mettransf.
DR Pfam; PF00590; TP_methylase; 2.
DR TIGRFAMS; TIGR01467; cobi chlL; 1.
DR TIGRFAMS; TIGR01466; cobiJ chlH; 1.
DR PROSITE; PS00839; SUMT 1; 1.
DR PROSITE; PS00840; SUMT 2; 1.
KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW Methyltransferase; Multifunctional enzyme; Complete proteome.
FT DOMAIN 1 243 PRECORRIN-2 C20-METHYLTRANSFERASE.
FT DOMAIN 244 508 PRECORRIN-3 METHYLASE.
SQ SEQUENCE 508 AA; 53910 MW; 95AC066F022C4DC1 CRC64;

Query Match
Best Local Similarity 56.0%; Score 42; DB 1; Length 508;
Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPGEQE 15
Db 247 GTVAVVGLGFGDSD 260

RESULT 4
RUVX CAUCR
ID RUVX CAUCR STANDARD; PRT; 156 AA.
AC Q9A5K8;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DE Putative Holliday junction resolvase (EC 3.1.1.-.-).
GN CC2439.
OS Caulobacter crescentus.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;
OC Caulobacteraceae; Caulobacter.
OX NCBI_TaxID=155892;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 39089 / CB15;
RX MEDLINE=21173698; PubMed=11259647;
RA Nierman W.C., Feldblyum T.V., Laub M.T., Paulsen I.T., Nelson K.E.,
RA Eisen J.A., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,
RA Potocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,
RA DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.B., Haft D.H.,
RA Kolonay J.F., Smit J., Craven M.B., Khouri H., Shetty J., Berry K.,
RA Utterback T., Tran K., Wolf A., Vamathevan J., Ermolaeva M., White O.,
RA Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;
RA "Complete genome sequence of Caulobacter crescentus.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001)
CC -!- FUNCTION: Could be a nuclease that resolves Holliday junction
CC intermediates in genetic recombination.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (Potential).
CC -!- SIMILARITY: Belongs to the YggF HJR family.
CC -----
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CC -----
CC EMBL; AE005913; AA024410.1; -.
CC PIR; F87551; F87551.
CC TIGR; CC2439; -.
CC HAMAP; MF 00651; -.
CC InterPro; IPR005227; Cons hypoth250.
CC Dr PIR; IPR006641; YggFC.
CC Pfam; PF03652; UPP0081; 1.

DR PIR; E70764; E70764.
DR TIGR; MT2126; -.
DR TubercuList; RV2066; -.
DR InterPro; IPR006364; Cobi ChlL.
DR InterPro; IPR006363; CobiJ.
DR InterPro; IPR008878; Cox/pox Metransf.
DR InterPro; IPR003043; Uropor_Mettransf.
DR Pfam; PF00590; TP_methylase; 2.
DR TIGRFAMS; TIGR01467; cobi chlL; 1.
DR TIGRFAMS; TIGR01466; cobiJ chlH; 1.
DR PROSITE; PS00839; SUMT 1; 1.
DR PROSITE; PS00840; SUMT 2; 1.
KW Cobalamin biosynthesis; Porphyrin biosynthesis; Transferase;
KW Methyltransferase; Multifunctional enzyme; Complete proteome.
FT DOMAIN 1 243 PRECORRIN-2 C20-METHYLTRANSFERASE.
FT DOMAIN 244 508 PRECORRIN-3 METHYLASE.
SQ SEQUENCE 508 AA; 53910 MW; 95AC066F022C4DC1 CRC64;

Query Match
Best Local Similarity 56.0%; Score 42; DB 1; Length 508;
Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPGEQE 15
Db 247 GTVAVVGLGFGDSD 260

RESULT 5
MDCG_XANAC
ID MDCG_XANAC STANDARD; PRT; 213 AA.
AC Q8PEW9;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Phosphoribosyl-dephospho-CoA transferase (EC 2.7.7.-) (Holo-ACP
DE synthase).
DE GN MDCG OR XAC0564.
OS Xanthomonas axonopodis (pv. citri).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=92829;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=306 / ATCC 13902 / XV 101;
RX MEDLINE=2202145; PubMed=12024217;
RA da Silva A.C.N., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
RA Camarotte G., Cannavan F., Cardoso J., Chamerigo F., Ciapina L.P.,
RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.,
RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
RA Pereira H.A.F., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
RA Spindola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
RA Setubal J.C., Kitajima J.P.;
RA "Comparison of the genomes of two Xanthomonas pathogens with differing
RA host specificities.";
RL Nature 417:459-463(2002)
CC -!- FUNCTION: Transfers 2-(5'-triphosphoribosyl)-3'-
CC dephosphocoenzyme-A to the apo-acyl carrier protein of the
CC malonate decarboxylase to yield holo-acyl carrier protein (By
CC similarity).
CC -!- CATALYTIC ACTIVITY: 2'-(5'-triphosphoribosyl)-3'-dephospho-CoA +
CC apo-ACP = holo-ACP + diphosphate.
CC -!- SIMILARITY: Belongs to the mdcG family.
CC -----
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CC -----
CC EMBL; AE011683; AA035453.1; -.
CC HAMAP; MF 00650; -.
CC Transferase; Nucleotidyltransferase; Complete proteome.
CC ACT_SITE 135 135 BY SIMILARITY.
CC ACT_SITE 137 137 BY SIMILARITY.
CC SEQUENCE 213 AA; 23013 MW; C378D8A975C75A17 CRC64;

```

Query Match 54.7%; Score 41; DB 1; Length 213;  
 Best Local Similarity 64.3%; Pred. No. 23;  
 Matches 9; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 AGLPAVUGLSPGE 14  
 DB 33 AGLPAVARGDSQ 46

RESULT 6  
 AT15 YEAST  
 ID AT15 YEAST STANDARD; PRT; 520 AA.  
 AC P25641; O8N1L6;  
 DT 01-MAY-1992 (Rel. 22, Created)  
 DT 15-MAR-2004 (Rel. 43, Last sequence update)  
 DE Autophagy related putative lipase ATG15 (EC 3.1.1.3) (Cytoplasm to vacuole targeting protein 17)  
 GN ATG15 OR AUT5 OR Cvt17 OR YCR068W OR YCR68W.  
 OS Saccharomyces cerevisiae (Baker's yeast).  
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;  
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.  
 OX NCBI\_TaxID=4932;  
 RN [1]  
 RP SEQUENCE FROM N.A., FUNCTION, GLYCOSYLATION, AND MUTAGENESIS OF SER-332.  
 RX MEDLINE=21125771; PubMed=11085977;  
 RA Teter S.A., Eggerton K.P., Scott S.V., Kim J., Fischer A.M., Klionsky D.J.,  
 RA "Degradation of lipid vesicles in the yeast vacuole requires function of Cvt17, a putative lipase.",  
 RL J. Biol. Chem. 276:2083-2087(2001).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=S288c;  
 RX MEDLINE=92243356; PubMed=1574125;  
 RA Oliver S.G., van der Aart Q.J.M., Agostoni-Carbone M.L., Aigle M., Alberghina L., Alexandraki D., Antoine G., Anwar R., Ballesta J.P.G., Benit P., Berben G., Bergantino E., Biteau N., Bolle P.-A., Botin-Fukuhara M., Brown A.J.P., Brown R., Buhler J.-M., Carignani G., Chanet R., Contreras R., Crouzet M., Daignan-Fornier B., De Haan M., Defoor E., Delgado M.D., Demolder J., Doira C., Dubois E., Dujon B., Dueterhoft A., Erdmann D., Esteban M., Fabre F., Fairhead C.A., Faye G., Feldmann H., Fiers W., Frangou-Gaillard M.-C., Franco L., Frontali L., Fukuhara H., Fuller L.J., Gent M.B., Gigot D., Gilliquet V., Glansdorff N., Goffeau A., Grenson M., Grisanti P., Grivelli L.A., Haasemann M., Hatat D., Hegemann J.H., Herbert C.J., Hilger F., Hohmann S., Hollenberg C.P., Huse K., Iborra F., Indge K.J., Isono K., Jackman P., Jacq C., Jacquet M., James C.M., Jauniaux J.-C., Jia Y., Jimenez A., Lucchini G., Lutzenkirchen K., Maat C., Mannhaupt G., Manzano M.E., Messenguy F., Mewes H.-W., Molemans F., McConnell D., McKee R.A., Nelon C.S., Olson M.V., Pallier C., Panzeri L., Pearson B.M., Perea J., Philippsen P., Pierard A., Planta R.J., Plevani P., Poetsch B., Pohl F.M., Purnelle B., Ramezani Rad M., Rasmussen S.W., Raynal A., Remacha M., Richterauer P., Roberts A.B., Rodriguez F., Sanz E., Schaaff-Gerstenschlaeger I., Scherens B., Schweitzer B., Shu Y., Skala J., Slonimski P.P., Sor F., Soustelle C., Spiegelberg R., Staveva L.I., Steensma H.Y., Steiner S., Thierry A., Threes G., Triano L.N., Urrestazu L.A., Valle G., Vetter I., van Vliet-Reedijk J.C., Volckaert G., Vreken P., Warrington J.R., von Wettstein D., Wicksteed B.L., Wilson C., Wurst H., Xu G., Zimmermann F.K., Sgouros J.G.;  
 RT "The complete DNA sequence of yeast chromosome III.";  
 RL Nature 357:38-46(1992).  
 RN [3]  
 EN REVISIONS TO C-TERMINUS.  
 RA Valles G., Volckaerts G.;  
 RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
 RN [4]  
 RP FUNCTION, SUBCELLULAR LOCATION, GLYCOSYLATION, AND MUTAGENESIS OF

RP SER-332.  
 RX MEDLINE=21450820; PubMed=11566994;  
 RA Epple U.D., Suriapranata I., Eskelinen E.-L., Thumm M.;  
 RT "Aut5/Cvt17p, a putative lipase essential for disintegration of autophagic bodies inside the vacuole.";  
 RL J. Bacteriol. 183:5942-5955(2001).  
 RN [5]  
 RP FUNCTION, SUBCELLULAR LOCATION, TOPOLOGY, AND GLYCOSYLATION.  
 RX MEDLINE=22499636; PubMed=12499386;  
 RA Epple U.D., Eskelinen E.-L., Thumm M.;  
 RT "Intravacuolar membrane lysis in Saccharomyces cerevisiae. Does vacuolar targeting of Cvt17/Aut5p affect its function?";  
 RL J. Biol. Chem. 278:7810-7821(2003).  
 CC -!- FUNCTION: Essential for lysis of subvacuolar cytoplasm to vacuole targeted bodies and intravacuolar autophagic bodies. Involved in the lysis of intravacuolar multivesicular body (MVB) vesicles.  
 CC -!- CATALYTIC ACTIVITY: Triacylglycerol + H(2)O = diacylglycerol + a fatty acid anion.  
 CC -!- SUBCELLULAR LOCATION: Type II membrane protein. Endoplasmic reticulum. From ER, targeted to vacuolar lumen at the MVB vesicles via the Golgi and the prevacuolar compartment (PVC).  
 CC -!- PTM: Glycosylated.  
 CC -!- SIMILARITY: Belongs to the AB hydrolase superfamily. Lipase family.  
 CC  
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 CC  
 DR EMBL; X59720; CAC42987.1; -.  
 DR PIR; S19483; S19483.  
 DR PIR; S74292; S74292.  
 DR GerOnline; 138970; -.  
 DR SGD; S0000664; ATG15.  
 DR GO; GO:0016021; C: integral to membrane; IPI.  
 DR GO; GO:0005775; C: vacuolar lumen; IDA.  
 DR GO; GO:0016298; F: lipase activity; IMP.  
 DR GO; GO:0006914; P: autophagy; IDA.  
 DR GO; GO:0030397; P: membrane degradation; IDA.  
 DR GO; GO:0006624; P: vacuolar protein processing/maturation; IDA.  
 DR InterPro; IPR002921; Lipase\_3.  
 DR InterPro; IPR008262; Lipase\_AS.  
 DR Pfam; PF01764; Lipase\_3; 1.  
 DR PROSITE; PS00120; LIPASE\_SER; 1.  
 DR Hydrolase; Lipid degradation; Autophagy; Endoplasmic reticulum; Signal-anchor; Transmembrane; Glycoprotein.  
 FT DOMAIN 1 14  
 FT TRANSMEM 15 35  
 FT DOMAIN 36 520  
 FT ACT\_SITE 332 332  
 FT CARBOHYD 173 173  
 FT CARBOHYD 202 202  
 FT CARBOHYD 208 208  
 FT MUTAGEN 332 332  
 SQ SEQUENCE 520 AA; 58435 MW; B56ABAB72B999019 CRC64;  
 Query Match 54.7%; Score 41; DB 1; Length 520;  
 Best Local Similarity 75.0%; Pred. No. 56;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVUGLSPGE 13  
 DB 346 GLPAVAFESPE 357

RESULT 7  
 LOL2\_HUMAN  
 ID LOL2\_HUMAN STANDARD; PRT; 774 AA.  
 AC Q9Y4K0; Q9BW70; Q9Y5Y8;

```

Query Match          54.7%; Score 41; DB 1; Length 774;
Best Local Similarity 66.7%; Pred. No. 83;
Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps

QY      2 GLPAVVGLSQCE 13
        |||||
Db      294 GLPAVWSCVPGQ 305

RESULT 8
RISB_PYRFU
AC Q8U4L8;
ID RISB_PYRFU STANDARD; PRT; 157 AA.
DT 10-OCT-2003 (Rel. 42, Created)
DI 10-OCT-2003 (Rel. 42, Last sequence update)
DD 10-OCT-2003 (Rel. 42, Last annotation update)
DE 6,7-dimethyl-8-ribitylumazine synthase (EC 2.5.1.9) (DMRL synthase)
DE (Lumazine synthase) (Riboflavin synthase beta chain).
DI RBH OR PF0063.
OS Pyrococcus furiosus.
OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;
OC Pyrococcus.
OC Pyrococcus.
OX NCBI_taxid=2261;
[1]
SEQUENCE FROM N.A.
STRAIN=Vc1 / DSM 3638 / ATCC 43587 / JCM 8422;
Weiss R.B., Dunn D.M., Robb F.T., Brown J.R.;
"the complete sequence of the Pyrococcus furiosus genome."
Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
!- FUNCTION: Riboflavin synthase is a bifunctional enzyme complex

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HSPSP; P00883; IADO.
DR InterPro; IPR000741; Aldolase I.
DR Pfam; PF00274; glycolytic enzy; 1.
DR ProDom; PD001128; Aldolase I; 1.
DR PROSITE; PS00158; ALDOLASE_CLASS_I; 1.
KW Lyase; Schiff base; Glycolysis; Multigene family.
FT BINDING 52 52 C-1-PHOSPHATE GROUP OF THE SUBSTRATE
  (BY SIMILARITY).
FT BINDING 143 143 C-1-PHOSPHATE GROUP OF THE SUBSTRATE
  (BY SIMILARITY).
FT BINDING 226 226 SCHIFF-BASE WITH DIHYDROXYACETONE-P
  (BY SIMILARITY).
FT ACT_SITE 359 359 ESSENTIAL FOR ENHANCED ACTIVITY OF THE
  ENZYME TOWARD FRUCTOSE 1,6-BISPHOSPHATE
  AS COMPARED WITH FRUCTOSE 1-PHOSPHATE
  (BY SIMILARITY).
SQ SEQUENCE 359 AA; 38490 MW; C0CAB16B9C1B9EF CRC64;

Query Match 53.3%; Score 40; DB 1; Length 359;
Best Local Similarity 60.08; Pred. No; 56;
Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps

Qy 1 AGLPAVVGLSPGQE 15
   | : | | | | | : |
Db 258 AAVPAVVFLSGGQSE 272

RESULT 10
ALF_CICAR
ID -ALF_CICAR STANDARD; PRT; 359 AA.
AC O65735;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Fructose-bisphosphate aldolase, cytoplasmic isozyme (EC 4.1.2.13).
GN ALDC.
OS Cicer arietinum (Chickpea) (Garbanzo).
OC Eukaryota; Viridiplantae; Streptophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids I; Fabales; Fabaceae; Papilionoideae; Cicereae; Cicer.
ON NCBI_TaxID=3827;
RX [1]
RN SEQUENCE FROM N.A.
RC STRAIN=cv. Castellana; TISSUE=Etiolated epicotyl;
RA Dopic B., Munoz F.J., Labrador E.;
RT "cDNA and deduced amino-acid sequence of a cytosolic aldolase from
RT Cicer arietinum L. epicotyls.";
RL (in) Plant Gene Register PGR98-110.
CC -1- CATALYTIC ACTIVITY: D-fructose 1,6-bisphosphate = glycerone
  phosphate + D-glyceraldehyde 3-phosphate.
CC -1- PATHWAY: Glycolysis; sixth step.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic.
CC -1- SIMILARITY: Belongs to class I fructose-bisphosphate aldolase
  family.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AJ005041; CAA06308.1; -.
DR HSPSP; P00883; IADO.
DR InterPro; IPR000741; Aldolase I.
DR Pfam; PF00274; glycolytic enzy; 1.
DR ProDom; PD001128; Aldolase I; 1.
DR PROSITE; PS00158; ALDOLASE_CLASS_I; 1.
KW Lyase; Schiff base; Glycolysis.
FT BINDING 52 52 C-1-PHOSPHATE GROUP OF THE SUBSTRATE.
FT BINDING 142 142 C-1-PHOSPHATE GROUP OF THE SUBSTRATE.
FT BINDING 226 226 SCHIFF-BASE WITH DIHYDROXYACETONE-P.

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FT ACT_SITE 359 359 ESSENTIAL FOR ENHANCED ACTIVITY OF THE
FT ENZYME TOWARD FRUCTOSE 1,6-BISPHOSPHATE
FT AS COMPARED WITH FRUCTOSE 1-PHOSPHATE.
SQ SEQUENCE 359 AA; 38451 MW; DD6864B745A5195 CRC64;

Query Match 53.3%; Score 40; DB 1; Length 359;
Best Local Similarity 60.0%; Pred. No. 56;
Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AGLPAVGLSPGEQ 15
Db 258 AAVPAVFLSGQSE 272

RESULT 11
VAL1_TM0V STANDARD; PRT; 361 AA.
ID VAL1_TM0V
AC Q06567;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE AL1 protein.
GN AL1.
OS Tomato mottle virus (isolate Florida) (TMoV).
OC Viruses; ssDNA viruses; Geminiviridae; Begonovirus.
OX NCBI_TaxID=36449;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=93107858; PubMed=1469361;
RA Abouzid A.M., Polston J.E., Hiebert E.;
RT "The nucleotide sequence of tomato mottle virus, a new geminivirus
RT isolated from tomatoes in Florida.";
RL J. Gen. Virol. 73:3225-3229 (1992).
CC -1- SIMILARITY: Belongs to the geminiviruses AL1 protein family.
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CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: L14460; AAC32414.1; --
CC PIR: JQ1870; JQ1870.
CC InterPro: IPR001191; Gemini_AL1.
CC Pfam: PF00799; Gemini_AL1; 1.
CC PRINTS: PR00227; GEMCOATL1.
CC ProDom: PD000736; Gemini_AL1; 1.
CC KW ATP-binding.
FT NP_BIND 222 229 ATP (BY SIMILARITY).
SQ SEQUENCE 361 AA; 40516 MW; 813B65CEBAC6950 CRC64;

Query Match 53.3%; Score 40; DB 1; Length 361;
Best Local Similarity 58.3%; Pred. No. 56;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 GLPAVGLSPGE 13
Db 299 GIPAVLCPNFE 310

RESULT 12
PLB5_SCHPO STANDARD; PRT; 633 AA.
ID PLB5_SCHPO
AC Q9Y7N6;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Putative lysophospholipase C1450.09c precursor (EC 3.1.1.5)
DE (Phospholipase B).
GN SPCC1450.09c.
OS Schizosaccharomyces pombe (Fission yeast).

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OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21849401; PubMed=11859360;
RA Wood V., Williams R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgouras J., Peat N., Hayes J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Fellwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford J., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volckaert G., Aert R., Robben J., Grymonprez B.,
RA Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Moeati D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shipakowski G.V., Ussery D., Barrell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe.";
RL Nature 415:871-880 (2002).
CC -1- FUNCTION: Catalyzes the release of fatty acids from
CC lysophospholipids (By similarity).
CC -1- CATALYTIC ACTIVITY: 2-lysophosphatidylcholine + H(2)O =
CC glycerophosphocholine + a fatty acid anion.
CC -1- SUBCELLULAR LOCATION: Secreted (Probable).
CC -1- SIMILARITY: Belongs to the lysophospholipase family.
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CC -----
CC EMBL: AL049559; CAB40176.2; --
CC GeneDB: SPombe; SPCC1450.09c; --
CC InterPro: IPR002642; PLAC.
CC Pfam: PF01735; PLA2_B; 1.
CC SMART: SM00022; PLAC; 1.
CC KW Hypothetical protein; Lipid degradation; Hydrolase; Glycoprotein;
CC Signal.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 633 PUTATIVE LYSOPHOSPHOLIPASE C1450.09C.
FT CARBOHYD 118 118 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 153 153 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 187 187 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 232 232 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 256 256 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 264 264 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 293 293 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 331 331 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 360 360 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 367 367 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 400 400 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 403 403 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 474 474 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 508 508 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 513 513 N-LINKED (GLCNAC. .) (POTENTIAL).

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FT CARBOHYD 537 537 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 564 564 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 586 586 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 603 603 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 SQ SEQUENCE 633 AA; 68292 MW; 49871B2955893D19 CRC64;  
 Query Match 53.3%; Score 40; DB 1; Length 633;  
 Best Local Similarity 75.0%; Pred. No. 97;  
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 PAVVGLSPGQE 15  
 DB 76 PASDGLSTGQE 87

RESULT 13  
 YHGF\_ECOLI STANDARD; PRT; 773 AA.  
 AC P46837; P76689;  
 DT 01-NOV-1995 (Rel. 32, Created)  
 DT 15-JUL-1999 (Rel. 38, Last sequence update)  
 DT 16-OCT-2001 (Rel. 40, Last annotation update)  
 DE Protein yHGF.  
 GN YHGF OR B3407.  
 OS Escherichia coli.  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;  
 OC Enterobacteriaceae; Escherichia.  
 OX NCBI\_TaxID=562;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=K12 / MG1655;  
 RX MEDLINE=97426617; PubMed=9278503;  
 RA Fountoulakis M., Takacs M.-P., Berndt P., Langen H., Takacs B.;  
 RA Biletner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,  
 RA Riley M., Collado-Vides J., Glaser J.D., Rode C.K., Mayhew G.F.,  
 RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,  
 RA Mau B., Shao Y.;  
 RT "The complete genome sequence of Escherichia coli K-12.";  
 RL Science 277:1453-1474(1997).  
 RN [2]  
 RP IDENTIFICATION BY MASS SPECTROMETRY.  
 RX MEDLINE=99420866; PubMed=10493123;  
 RA Fountoulakis M., Takacs M.-P., Berndt P., Langen H., Takacs B.;  
 RT "Enrichment of low abundance proteins of Escherichia coli by  
 RT hydroxyapatite chromatography";  
 RL Electrophoresis 20:2181-2195(1999).  
 CC -!- SIMILARITY: STRONG, TO H.INFLUENZAE HI0568.  
 CC -!- SIMILARITY: Contains 1 SI motif domain.  
 CC  
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 CC  
 CC EMBL; U18997; AAA58204.1; ALT\_FRAME.  
 DR EMBL; U18997; AAA58205.1; ALT\_FRAME.  
 DR EMBL; AE000416; AAC76432.1; ALT\_INIT.  
 DR HSP; P05055; ISRO.  
 DR Ecogen; EGI2932; yHGF.  
 DR InterPro; IPR008994; Nucleic\_acid\_OB.  
 DR InterPro; IPR003029; S1.  
 DR InterPro; IPR006641; YqgFc.  
 DR Pfam; PF00575; S1; 1.  
 DR SMART; SM00316; S1; 1.  
 DR SMART; SM00732; YqgFc; 1.  
 DR PROSITE; PS0126; S1; 1.  
 KW RNA-binding; Complete proteome.  
 FT DOMAIN 651 720 SI MOTIF.  
 FT CONFLICT 754 755 QP -> HA (IN REF. 1; AAA58205).  
 SQ SEQUENCE 773 AA; 85119 MW; EA54D9ED952A8229 CRC64;

QY 1 AGUPAVVGLSPG 12  
 DB 324 AGLRATMGIDPG 335

Query Match 53.3%; Score 40; DB 1; Length 773;  
 Best Local Similarity 66.7%; Pred. No. 1.2e+02;  
 Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGUPAVVGLSPG 12  
 DB 324 AGLRATMGIDPG 335

RESULT 14  
 CADM\_MOUSE STANDARD; PRT; 813 AA.  
 AC Q9WTE5;  
 DT 28-FEB-2003 (Rel. 41, Created)  
 DT 28-FEB-2003 (Rel. 41, Last sequence update)  
 DT 28-FEB-2003 (Rel. 41, Last annotation update)  
 DE Cadherin-22 precursor (PB-cadherin).  
 GN CDH22.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_TaxID=10090;  
 RN [1]  
 RP SEQUENCE FROM N.A., AND DEVELOPMENTAL STAGE.  
 RC STRAIN=ICR; TISSUE=Brain;  
 RX MEDLINE=99326347; PubMed=10398531;  
 RA Kitajima K., Koshimizu U., Nakamura T.;  
 RT "Expression of a novel type of classic cadherin, PB-cadherin in  
 RT developing brain and limb buds.";  
 RL Dev. Dyn. 215:206-214(1999).  
 CC -!- FUNCTION: Cadherins are calcium dependent cell adhesion proteins.  
 CC They preferentially interact with themselves in a homophilic  
 CC manner in connecting cells; cadherins may thus contribute to the  
 CC sorting of heterogeneous cell types. PB-cadherins may have a role  
 CC in the morphological organization of pituitary gland and brain  
 CC tissues  
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein.  
 CC -!- TISSUE SPECIFICITY: Predominantly expressed in brain. Abundant in  
 CC olfactory bulb, cerebrum, and cerebellum, less in pons, medulla,  
 CC and spinal cord. Low expression in heart. No expression in lung,  
 CC liver, spleen, kidney, testis, stomach, intestine, colon, and  
 CC placenta.  
 CC -!- DEVELOPMENTAL STAGE: Expressed at 9.5 dpc onwards. At 10.5 dpc, in  
 CC brain (telencephalic vesicles and isthmus), spinal cord and limb  
 CC buds (in the zone of polarizing activity). At 14.5 dpc, in  
 CC olfactory bulb and cerebellum.  
 CC -!- INDUCTION: Down-regulated by thyroid hormone.  
 CC -!- SIMILARITY: Contains 5 cadherin domains.  
 CC  
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 CC  
 CC EMBL; AB019618; BAA34426.1; -.  
 DR EMBL; MGI:1341843; Cdh22.  
 DR HSSP; P15116; INCU.  
 DR InterPro; IPR002126; Cadherin.  
 DR InterPro; IPR002333; Cadherin\_C\_term.  
 DR Pfam; PF00028; cadherin; 5.  
 DR Pfam; PF01049; Cadherin\_C\_term; 1.  
 DR PRINTS; PR00205; CADHERIN.  
 DR SMART; SM00112; CA; 5.  
 DR PROSITE; PS00232; CADHERIN\_1; 2.  
 DR PROSITE; PS0268; CADHERIN\_2; 5.  
 KW Cell adhesion; Glycoprotein; Transmembrane; Calcium-binding; Repeat;  
 KW SIGNAL.  
 FT SIGNAL 1 33 POTENTIAL.  
 FT CHAIN 34 813 CADHERIN-22.  
 FT DOMAIN 33 621 EXTRACELLULAR (POTENTIAL).

FT TRANSMEM 622 642 POTENTIAL.  
 FT DOMAIN 643 813 CYTOPLASMIC (POTENTIAL).  
 FT DOMAIN 61 165 CADHERIN 1.  
 FT DOMAIN 166 274 CADHERIN 2.  
 FT DOMAIN 275 391 CADHERIN 3.  
 FT DOMAIN 392 495 CADHERIN 4.  
 FT DOMAIN 496 613 CADHERIN 5.  
 FT CARBOHYD 159 159 N-LINKED (GLCNAC. .) (POTENTIAL).  
 FT CARBOHYD 463 463 N-LINKED (GLCNAC. .) (POTENTIAL).  
 FT CARBOHYD 609 609 N-LINKED (GLCNAC. .) (POTENTIAL).  
 SQ SEQUENCE 813 AA; 86021 MW; 5510F9848D976567 CRC64;  
 Query Match 53.3%; Score 40; DB 1; Length 813;  
 Best Local Similarity 60.0%; Pred. No. 1.2e+02;  
 Matches 9; Conservative 0; Mismatches 6; Indels 0; Gaps 0;  
 QY 1 AGLPAVVGLSPEQEF 15  
 DB 34 ASTPAPSSLSPGAQE 48  
 RESULT 15  
 PGCA BOVIN  
 ID PGCA BOVIN STANDARD; PRT; 2364 AA.  
 AC P13608; P79117; Q28159;  
 DT 01-JAN-1990 (Rel. 13, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 15-MAR-2004 (Rel. 43, Last annotation update)  
 DE Aggrecan core protein precursor (Cartilage-specific proteoglycan core protein) (CSCP).  
 DE protein) (CSCP).  
 GN AGCL.  
 OS Bos taurus (Bovine).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 OC Bovidae; Bovinae; Bos.  
 OC NCBI\_TaxID=9913;  
 RN [1]  
 RP Hering T.M., Kollar J., Huynh T.D.;  
 RA Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RP SEQUENCE OF 563-1056 FROM N.A.  
 RX MEDLINE=89380219; PubMed=2528543;  
 RA Antonsson P., Heinegaard D., Oldberg A.;  
 RT "The keratan sulfate-enriched region of bovine cartilage proteoglycan consists of a consecutively repeated hexapeptide motif.";  
 RN J. Biol. Chem. 264:16170-16173(1989).  
 RN [3]  
 RP SEQUENCE OF 1609-2113 AND 2151-2364 FROM N.A.  
 RX MEDLINE=87270630; PubMed=3111460;  
 RA Oldberg A., Antonsson P., Heinegaard D.;  
 RT "The partial amino acid sequence of bovine cartilage proteoglycan, deduced from a cDNA clone, contains numerous Ser-Gly sequences arranged in homologous repeats.";  
 RN Biochem. J. 243:255-259(1987).  
 RN [4]  
 RP SEQUENCE OF 2114-2150 FROM N.A.  
 RC TISSUE=Cartilage;  
 RX MEDLINE=93352525; PubMed=8349621;  
 RA Fuellep C., Waicz E., Vailon M., Glant T.T.;  
 RT "Expression of alternatively spliced epidermal growth factor-like domains in aggrecans of different species. Evidence for a novel module.";  
 RN J. Biol. Chem. 268:17377-17383(1993).  
 RN [5]  
 RP PARTIAL SEQUENCE.  
 RX MEDLINE=85027710; PubMed=6489519;  
 RA Perin J.-P., Bonnet F., Jolles J., Jolles P.;  
 RT "Sequence data concerning the protein core of the cartilage proteoglycan monomers. Characterization of a sequence allowing the synthesis of an oligonucleotide probe.";  
 RN FEBS Lett. 176:37-42(1984).  
 RN [6]

RP PARTIAL SEQUENCE.  
 RX MEDLINE=87005253; PubMed=3530809;  
 RA Perin J.-P., Bonnet F., Jolles P.;  
 RT "Structural relationship between link proteins and proteoglycan monomers.";  
 RN FEBS Lett. 206:73-77(1986).  
 CC -!- FUNCTION: This proteoglycan is a major component of extracellular matrix of cartilaginous tissues. A major function of this protein is to resist compression in cartilage. It binds avidly to hyaluronic acid via an amino-terminal globular region. May play a regulatory role in the matrix assembly of the cartilage.  
 CC -!- SUBCELLULAR LOCATION: Secreted; extracellular matrix (By similarity).  
 CC -!- ALTERNATIVE PRODUCTS:  
 CC Event=Alternative splicing; Named isoforms=2;  
 CC Name=1;  
 CC IsoId=P13608-1; Sequence=Displayed;  
 CC Name=2;  
 CC IsoId=P13608-2; Sequence=VSP\_003072;  
 CC -!- DOMAIN: Two globular domains, G1 and G2, comprise the amino terminus of the proteoglycan, while another globular region, G3, makes up the C-terminus. G1 contains link domains and thus consists of three disulfide-bonded loop structures designated as the A, B, B' motifs. G2 is similar to G1. The keratan sulfate (KS) and the chondroitin sulfate (CS) attachment domains lie between G2 and G3.  
 CC -!- PTM: CONTAINS MOSTLY CHONDROITIN SULFATE, BUT ALSO N-LINKED AND O-LINKED (ABOUT 40) OLIGOSACCHARIDES.  
 CC -!- PTM: THE KERATAN SULFATE CONTENTS DIFFER CONSIDERABLY BETWEEN ADULT AND FETAL BOVINE PROTEOGLYCANS.  
 CC -!- SIMILARITY: Contains 1 immunoglobulin-like V-type domain.  
 CC -!- SIMILARITY: Contains 4 link domains.  
 CC -!- SIMILARITY: Contains 1 EGF-like domain.  
 CC -!- SIMILARITY: Contains 1 C-type lectin family domain.  
 CC -!- SIMILARITY: Contains 1 Sushi (SCR) domain.  
 CC -!- SIMILARITY: Belongs to the aggrecan/versican proteoglycan family.  
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 -----  
 DR EMBL; U76615; AAB38524.1; -;  
 DR EMBL; L07053; -; NOT\_ANNOTATED\_CDS.  
 DR PIR; A34234; A39808.  
 DR PIR; T42630; T42630.  
 DR HSPSP; P08709; IBF9.  
 DR InterPro; IPR002353; Antifreeze1.  
 DR InterPro; IPR000152; Asx\_hydroxyl\_s.  
 DR InterPro; IPR000742; EGF-2.  
 DR InterPro; IPR001881; EGF\_Ca.  
 DR InterPro; IPR006209; EGF-like.  
 DR InterPro; IPR007110; Ig-Like.  
 DR InterPro; IPR003006; Ig\_MHC.  
 DR InterPro; IPR001304; Lectin\_C.  
 DR InterPro; IPR000538; Link.  
 DR InterPro; IPR003324; SGXSG.  
 DR InterPro; IPR000436; Sushi\_SCR\_CCP.  
 DR Pfam; PF00008; EGF; 1.  
 DR Pfam; PF00047; Ig; 1.  
 DR Pfam; PF00059; Lectin\_c; 1.  
 DR Pfam; PF02339; SGXSG; 61.  
 DR Pfam; PF00084; sushi; 1.  
 DR Pfam; PF00193; Xlink; 4.  
 DR PRINTS; PR00356; ANTIFREEZE1.  
 DR PRINTS; PR01285; LINKMODULE.  
 DR ProDom; PD000918; Link; 4.  
 DR SMART; SM00032; CCP; 1.  
 DR SMART; SM00034; CLECT; 1.  
 DR SMART; SM00179; EGF\_CA; 1.

DR SMART; SM00445; LINK; 4.  
 DR PROSITE; PS00010; ASX HYDROXYL; 1.  
 DR PROSITE; PS00615; C-TYPE LECTIN 1; 1.  
 DR PROSITE; PS0041; C-TYPE LECTIN 2; 1.  
 DR PROSITE; PS00022; EGF 1; 1.  
 DR PROSITE; PS00026; EGF 3; 1.  
 DR PROSITE; PS01187; EGF CA; 1.  
 DR PROSITE; PS00835; IG LIKE; 1.  
 DR PROSITE; PS00290; IG-MHC; FALSE\_NEG.  
 DR PROSITE; PS01241; LINK; 4.  
 KW Glycoprotein; Proteoglycan; Lectin; Signal; Sushi; EGF-like domain;  
 KW Calcium; Alternative splicing; Repeat; Immunoglobulin domain.  
 FT SIGNAL 1 16  
 FT CHAIN 17 2364  
 FT DOMAIN 25 147  
 FT DOMAIN 170 247  
 FT DOMAIN 268 349  
 FT DOMAIN 504 581  
 FT DOMAIN 602 683  
 FT DOMAIN 774 907  
 FT DOMAIN 1433 2112  
 FT DOMAIN 2113 2149  
 FT DOMAIN 2114 2364  
 FT DOMAIN 2161 2276  
 FT DOMAIN 2280 2338  
 FT DISULFID 51 133  
 FT DISULFID 175 246  
 FT DISULFID 199 220  
 FT DISULFID 273 348  
 FT DISULFID 297 318  
 FT DISULFID 509 580  
 FT DISULFID 533 554  
 FT DISULFID 607 682  
 FT DISULFID 631 652  
 FT DISULFID 2117 2128  
 FT DISULFID 2182 2274  
 FT DISULFID 2250 2266  
 FT DISULFID 2281 2324  
 FT DISULFID 2310 2337  
 FT CARBOHYD 126 126  
 FT CARBOHYD 239 239  
 FT CARBOHYD 333 333  
 FT CARBOHYD 387 387  
 FT CARBOHYD 611 611  
 FT CARBOHYD 667 667  
 FT VARSPLIC 2114 2150  
 SQ SEQUENCE 2364 AA; 246359 MW; 6FF83763420C3D4C CRC64;  
 Query Match 53.3%; Score 40; DB 1; Length 2364;  
 Best Local Similarity 53.3%; Pred. No. 3.6e+02;  
 Matches 8; Conservative 2; Mismatches 5; Indels 0; Gaps 0;  
 QY 1 AGLPAVGLSPGEQE 15  
 Db :||| |||  
 971 SGLPVEGLSPSGEE 985

Search completed: May 7, 2004, 12:34:34  
 Job time : 6.4 secs

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OM protein - protein search, using sw model

Run on: May 7, 2004, 12:26:40 ; Search time 28.05 Seconds  
(without alignments)  
168.726 Million cell updates/sec

Title: US-09-786-214A-15

Perfect score: 75

Sequence: 1 AGLPAVVLSPGEQE 15

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:\*

- 1: sp\_archea:\*
- 2: sp\_bacteria:\*
- 3: sp\_fungi:\*
- 4: sp\_human:\*
- 5: sp\_invertebrate:\*
- 6: sp\_mammal:\*
- 7: sp\_mhc:\*
- 8: sp\_organelle:\*
- 9: sp\_phage:\*
- 10: sp\_plant:\*
- 11: sp\_rodent:\*
- 12: sp\_virus:\*
- 13: sp\_vertebrate:\*
- 14: sp\_unclassified:\*
- 15: sp\_rvirus:\*
- 16: sp\_bacteriap:\*
- 17: sp\_archeap:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	55	73.3	41	11 Q8K408	Q8K408 rattus norv
2	50	66.7	381	16 Q9RL15	Q9RL15 deinococcus
3	46	61.3	150	16 Q8PE55	Q8PE55 xanthomonas
4	46	61.3	422	16 Q9KZB0	Q9KZB0 streptomyces
5	44	58.7	335	16 Q8ZRR7	Q8ZRR7 streptomyces
6	44	58.7	383	16 Q7W8U3	Q7W8U3 prochlorococcus
7	44	58.7	399	16 Q8DKG5	Q8DKG5 synecococcus
8	43.5	58.0	614	16 Q8Y280	Q8Y280 ralstonia s
9	43	57.3	579	16 Q8NEJ7	Q8NEJ7 corynebacte
10	43	57.3	582	16 Q8FQM6	Q8FQM6 corynebacte
11	43	57.3	821	17 Q9HPR8	Q9HPR8 halobacteri
12	42.5	56.7	305	16 Q89FB7	Q89FB7 bradyrhizob
13	42	56.0	254	16 Q89BH1	Q89BH1 bradyrhizob
14	42	56.0	326	1 Q9UXP0	Q9UXP0 methanobact
15	42	56.0	574	2 Q54756	Q54756 synecococcus
16	42	56.0	660	9 Q7Y4W9	Q7Y4W9 bacterioph

17	42	56.0	673	13 Q7T2X8	Q7T2X8 gallus gall
18	42	56.0	1541	6 Q8HXL3	Q8HXL3 sus scrofa
19	41	54.7	143	4 Q8NAW2	Q8NAW2 homo sapien
20	41	54.7	164	16 Q99QD9	Q99QD9 caulobacter
21	41	54.7	177	17 Q9YAM1	Q9YAM1 aeropyrum p
22	41	54.7	214	16 Q9I2S0	Q9I2S0 pseudomonas
23	41	54.7	222	17 Q978I4	Q978I4 thermoplasma
24	41	54.7	239	16 Q89JT2	Q89JT2 bradyrhizob
25	41	54.7	438	17 Q8QOK1	Q8QOK1 methanobact
26	41	54.7	448	16 Q8PML2	Q8PML2 xanthomonas
27	41	54.7	520	3 Q8N1L6	Q8N1L6 saccharomyc
28	41	54.7	529	4 Q8N1S8	Q8N1S8 homo sapien
29	41	54.7	11096	2 Q9L4W3	Q9L4W3 streptomyces
30	40	53.3	85	6 Q7786	Q7786 canis famill
31	40	53.3	130	17 Q9YFJ5	Q9YFJ5 aeropyrum p
32	40	53.3	154	17 Q8ZJX3	Q8ZJX3 pyrobaculum
33	40	53.3	242	16 Q7WCX1	Q7WCX1 bordetella
34	40	53.3	261	16 Q8NOA2	Q8NOA2 corynebacte
35	40	53.3	278	2 Q9F0Y4	Q9F0Y4 xanthomonas
36	40	53.3	298	16 Q7W5D6	Q7W5D6 bordetella
37	40	53.3	298	16 Q7VII2	Q7VII2 bordetella
38	40	53.3	331	16 Q7WEG7	Q7WEG7 bordetella
39	40	53.3	331	16 Q7W350	Q7W350 bordetella
40	40	53.3	331	16 Q7W032	Q7W032 bordetella
41	40	53.3	355	10 Q947A7	Q947A7 nitellopsis
42	40	53.3	357	4 Q8NEX1	Q8NEX1 homo sapien
43	40	53.3	358	4 Q9NWD0	Q9NWD0 homo sapien
44	40	53.3	358	10 Q9FUG7	Q9FUG7 fragaria an
45	40	53.3	358	12 Q91201	Q91201 havana toma

## ALIGNMENTS

RESULT 1

Q8K408 PRELIMINARY; PRT; 41 AA.

AC Q8K408; 01-OCT-2002 (TREMBLrel. 22, Created)

DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)

DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)

DE Truncated macrophage colony stimulating factor.

GN CSF1.

OS Rattus norvegicus (Rat).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OX NCBI\_TaxID=10116;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=LEW.tl;

RX MEDLINE=22069908; PubMed=12074592;

RA Dobbins D.E.; Sood R., Hashimoto A., Hansen C.T., Wilder R.L.,

RA Remmers E.F.;

RT "Mutation of macrophage colony stimulating factor (CSF1) causes osteopetrosis in the tl rat."

RL Biochem. Biophys. Res. Commun. 294:1114-1120(2002).

DR EMBL; AF514357; AAM54137.1; -

SQ SEQUENCE 41 AA; 4178 MW; ID342C19BD18AA41 CRC64;

Query Match 73.3%; Score 55; DB 11; Length 41;  
Best Local Similarity 78.6%; Pred. No. 0.1;  
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVLSPGEQE 15  
Db 20 GLPAVVLSPGEQE 33

RESULT 2

Q9RL15 PRELIMINARY; PRT; 381 AA.

ID Q9RL15; 01-MAY-2000 (TREMBLrel. 13, Created)

DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)  
 DE 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Cytochrome P450.  
 GN DR2473.  
 OS Deinococcus radiodurans.  
 OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;  
 OC Deinococcaceae; Deinococcus.  
 OX NCBI\_TaxID=1299;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=RL / ATCC 13939 / DSM 20539 / NCIB 9279;  
 RX MEDLINE=20036896; PubMed=10567266;  
 RA White O., Eissen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,  
 RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,  
 RA Moffat K.S., Qin H., Jiang L., Pauphille W., Crosby M., Shen M.,  
 RA Vamathevan J.J., Lam P., McDonald J., Utterback T., Zalewski C.,  
 RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,  
 RA Ketchum K.A., Aravind K.E., Salzberg S., Smith H.O., Venter J.C.,  
 RA Fraser C.M.;  
 RT "Genome sequence of the radioresistant bacterium Deinococcus  
 RL radiodurans R1.";  
 RL Science 286:1571-1577(1999)  
 CC -!- SIMILARITY: BELONGS TO THE CYTOCHROME P450 FAMILY.  
 DR EMBL; AE002076; AAF12016.1; -.  
 DR FIR; F75270; F75270.  
 DR TIGR; DR2473; -.  
 DR GO; GO:0004497; F:monooxygenase activity; IEA.  
 DR GO; GO:0006118; P:electron transport; IEA.  
 DR InterPro; IPR001128; Cytochrome\_P450.  
 DR Pfam; PF00067; P450; 1.  
 DR PRINTS; PR00385; P450.  
 DR PROSITE; PS00086; CYTOCHROME P450; 1.  
 DR Heme; Monooxygenase; Oxidoreductase; Complete proteome.  
 SQ SEQUENCE 381 AA; 41940 MW; F191EA69F1797B53 CRC64;  
  
 Query Match 66.7%; Score 50; DB 16; Length 381;  
 Best Local Similarity 100.0%; Pred. No. 7.4;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
 QY 2 GLPAVVGSLSP 11  
 DB 51 GLPAVVGSLSP 60  
  
 RESULT 3  
 Q8PE55 PRELIMINARY; PRT; 150 AA.  
 AC Q8PE55;  
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)  
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)  
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)  
 DE Deoxycytidylate deaminase.  
 GN XCC0126.  
 OS Xanthomonas campestris (pv. campestris).  
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;  
 OC Xanthomonadaceae; Xanthomonas.  
 OX NCBI\_TaxID=340;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 33913 / NCPPB 528;  
 RX MEDLINE=2022145; PubMed=12024217;  
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,  
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,  
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,  
 RA Camarotte G., Cannavan F., Cardoso J., Chambergo F., Ciapina L.P.,  
 RA Cicarelli R.M.B., Coutinho L.B., Cursino-Santos J.R., El-Dorri H.,  
 RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,  
 RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,  
 RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,  
 RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.H.,  
 RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,  
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,  
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,

RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,  
 RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,  
 RA Setubal J.C., Kitajima J.P.;  
 RT "Comparison of the genomes of two Xanthomonas pathogens with differing  
 RT host specificities.";  
 RL Nature 417:459-463(2002).  
 DR EMBL; AE012107; AAM39445.1; -.  
 DR GO; GO:0016787; F:hydrolase activity; IEA.  
 DR GO; GO:0008270; F:zinc ion binding; IEA.  
 DR InterPro; IPR002125; dCMP/cyt\_deam.  
 DR Pfam; PF00383; dCMP\_cyt\_deam; 1.  
 KW Complete proteome.  
 SQ SEQUENCE 150 AA; 16201 MW; 63D9C17D44DC9B43 CRC64;  
  
 Query Match 61.3%; Score 46; DB 16; Length 150;  
 Best Local Similarity 60.0%; Pred. No. 12;  
 Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;  
  
 QY 1 AGLPAVVGSLSPGEQ 15  
 DB 104 AGIKRWVALAFGESE 118  
  
 RESULT 4  
 Q3KZB0 PRELIMINARY; PRT; 462 AA.  
 ID Q3KZB0  
 AC Q3KZB0;  
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)  
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)  
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)  
 DE Putative gntR-family regulator.  
 GN SC04836 OR SC5G8.04.  
 OS Streptomyces coelicolor.  
 OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;  
 OC Streptomycineae; Streptomycetaceae; Streptomyces.  
 OX NCBI\_TaxID=1302;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=A3(2);  
 RA Saunders D.C., Harris D.;  
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=A3(2);  
 RA Bentley S.D., Parkhill J., Barrell B.G., Rajandream M.A.;  
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=A3(2);  
 RX MEDLINE=97000351; PubMed=8843436;  
 RA Redenbach M., Kieser H.M., Denapante D., Eichner A., Cullum J.,  
 RA Kinashi H., Hopwood D.A.;  
 RT "A set of ordered cosmids and a detailed genetic and physical map for  
 RT the 8 Mb Streptomyces coelicolor A3(2) chromosome.";  
 RL Mol. Microbiol. 21:77-96(1996).  
 RN [4]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=A3(2) / M145;  
 RX MEDLINE=21996410; PubMed=12000953;  
 RA Bentley S.D., Chater K.F., Cerdeno-Tarraga A.-M., Challis G.L.,  
 RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,  
 RA Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,  
 RA Cronin A., Fraser A., Gobie A., Hidalgo J., Hornsby T., Howarth S.,  
 RA Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neill S.,  
 RA Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,  
 RA Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor K.,  
 RA Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,  
 RA Hopwood D.A.;  
 RT "Complete genome sequence of the model actinomycete Streptomyces  
 RT coelicolor A3(2).";  
 RL Nature 417:141-147(2002).  
 CC -!- SIMILARITY: BELONGS TO THE GNTR FAMILY OF TRANSCRIPTIONAL  
 CC REGULATORS.

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DR EMBL; AL939121; CAB89055.1; -.
DR GO; GO:0005622; C:intracellular; IEA.
DR GO; GO:0003700; P:transcription factor activity; IEA.
DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR000524; HTH GntR.
DR InterPro; IPR000408; Reg_chir_condens.
DR Pfam; PF00392; gntR; 1.
DR PRINTS; PR00035; HTHGNTR.
DR SMART; SM00345; HTH GntR; 1.
DR PROSITE; PS00626; RCCL 2; 1.
KW DNA-binding; Transcription regulation; Complete proteome.
SQ SEQUENCE 462 AA; 48831 MW; DFCECA65LEI13317 CRC64;

Query Match 61.3%; Score 46; DB 16; Length 462;
Best Local Similarity 60.0%; Pred. No. 41;
Matches 9; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGLPAVVGLSPGEQ 15
DB 392 AGLHAVLGLPPGTEQ 406

RESULT 5
Q82RR7 PRELIMINARY; PRT; 335 AA.
AC Q82RR7;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Putative terpene cyclase.
GN TPC1 OR SAV76.
OS Streptomyces avermitilis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OC NCBI_TaxID=33903;
RN [1]
RC SEQUENCE FROM N.A.
RX STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=21477403; PubMed=11572948;
RA Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
RA Shinose M., Takahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
RA Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
RT "Genome sequence of an industrial microorganism Streptomyces
RT avermitilis: deducing the ability of producing secondary
RT metabolites."
RL Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=22608306; PubMed=12692562;
RA Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
RA Sakaki Y., Hattori M., Omura S.;
RT "Complete genome sequence and comparative analysis of the industrial
RT microorganism Streptomyces avermitilis."
RL Nat. Biotechnol. 21:526-531(2003).
RL EMBL; AP005021; BAC67785.1; -.
DR InterPro; IPR008949; Terpenoid_synth.
DR Complete proteome.
KW SEQUENCE 335 AA; 36480 MW; 49B8477E2D52666F CRC64;

Query Match 58.7%; Score 44; DB 16; Length 335;
Best Local Similarity 72.7%; Pred. No. 62;
Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPG 12
DB 8 GLPAPAGISPG 18

RESULT 6
Q7V8U3 PRELIMINARY; PRT; 383 AA.
ID Q7V8U3
AC Q7V8U3;

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DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Phage integrase.
GN PMT0234.
OS Prochlorococcus marinus (strain MIT 9313).
OC Bacteria; Cyanobacteria; Prochlorophytes; Prochlorococaceae;
OC Prochlorococcus.
OX NCBI_TaxID=74547;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22825698; PubMed=12917642;
RA Rocap G., Larimer F.W., Lamerdin J., Malfatti S., Chain P.,
RA Algren N.A., Arellano A., Coleman M., Hauser L., Hess W.R.,
RA Johnson Z.I., Land M., Lindell D., Post A.F., Regala W., Shah M.,
RA Shaw S.L., Steglich C., Sullivan M.B., Ting C.S., Tolonen A.,
RA Webb E.A., Zinser E.R., Chisholm S.W.;
RT "Genome divergence in two Prochlorococcus ecotypes reflects oceanic
RT niche differentiation."
RL Nature 424:1042-1047(2003).
DR EMBL; BX572095; CAE20409.1; -.
KW Complete proteome.
SQ SEQUENCE 383 AA; 43234 MW; CF956FE676F2E59B CRC64;

Query Match 58.7%; Score 44; DB 16; Length 383;
Best Local Similarity 75.0%; Pred. No. 71;
Matches 9; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPGE 13
DB 193 GLPASVRLTPGE 204

RESULT 7
Q8DKG5 PRELIMINARY; PRT; 399 AA.
AC Q8DKG5;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Tlr0894 protein.
GN TLR0894.
OS Synechococcus elongatus (Thermosynechococcus elongatus).
OC Bacteria; Cyanobacteria; Chroococcales; Synechococcus.
OC NCBI_TaxID=32046;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BP-1;
RX MEDLINE=22225144; PubMed=12240834;
RA Nakamura Y., Kaneko T., Sato S., Ikeuchi M., Kato H., Sasamoto S.,
RA Watanabe A., Iriguchi M., Kawashima K., Kimura T., Kishida Y.,
RA Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Nakazaki N.,
RA Shimpo S., Sugimoto M., Takeuchi C., Yamada M., Tabata S.;
RT "Complete genome structure of the thermophilic cyanobacterium
RT Thermosynechococcus elongatus BP-1."
RL DNA Res. 9:123-130(2002).
DR EMBL; AP005372; BAC08446.1; -.
DR InterPro; IPR000437; Prok_lipoprot_S.
DR PROSITE; PS00013; PROKAR_LIPOPROTEIN; 1.
KW Complete proteome.
SQ SEQUENCE 399 AA; 45259 MW; 61F60CE5B22A3F94 CRC64;

Query Match 58.7%; Score 44; DB 16; Length 399;
Best Local Similarity 64.3%; Pred. No. 74;
Matches 9; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVGLSPGEQ 15
DB 273 GLPFFVKGVSFYEQQ 286

RESULT 8
Q8Y280

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ID OBY280 PRELIMINARY; PRT; 614 AA.
AC OBY280;
DT 01-MAR-2002 (TREMBlrel. 20, Created)
DT 01-MAR-2002 (TREMBlrel. 20, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Probable ATP-binding transport ABC transporter protein.
GN RSC0456 OR RS0444.
OS Ralstonia solanacearum (Pseudomonas solanacearum).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Burkholderiaceae; Ralstonia.
OX NCBI_TaxID=305;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=GM11000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
RA Arlat M., Billault A., Brottier P., Camus J.C., Cattolico L.,
RA Chandler M., Choine N., Claudel-Renard C., Cunnac S., Demange N.,
RA Gaspin C., Lavie M., Moisan A., Robert C., Saurin W., Schiex T.,
RA Sigvier P., Thebault P., Whalen M., Wincker P., Levy M.,
RA Weisenbach J., Boucher C.A.;
RT "Genome sequence of the plant pathogen Ralstonia solanacearum.";
RL Nature 415:497-502(2002).
DR EMBL; AL646059; CAD13984.1; -.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0004009; F:ATP-binding cassette (ABC) transporter acti...; IEA.
DR GO; GO:0000166; F:nucleotide binding; IEA.
DR GO; GO:0006810; P:transport; IEA.
DR InterPro; IPR003593; AAA ATPase.
DR InterPro; IPR003439; ABC transporter.
DR InterPro; IPR001623; DnaJ N.
DR Pfam; PF00005; ABC tran. I.
DR ProDom; PD000006; ABC_transporter; 1.
DR SMART; SM00382; AAA; 1.
DR PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
DR PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
DR PROSITE; PS00636; DnaJ_1; 1.
KW Complete proteome.
SQ SEQUENCE 614 AA; 59240 MW; E293355B85872142 CRC64;

Query Match 58.0%; Score 43.5; DB 16; Length 614;
Best Local Similarity 45.8%; Pred. No. 1.4e+02;
Matches 11; Conservative 2; Mismatches 0; Indels 11; Gaps 1;

QY 3 LPAVVG-----LSPGQE 15
Db 501 LPALVGLDEVSNWLSRLSPGQQ 524

RESULT 9
Q8NRJ7 PRELIMINARY; PRT; 579 AA.
AC Q8NRJ7;
DT 01-OCT-2002 (TREMBlrel. 22, Created)
DT 01-OCT-2002 (TREMBlrel. 22, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Sulfate permease and related transporters (MFS superfamily).
GN CGL1051.
OS Corynebacterium glutamicum (Brevibacterium flavum).
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Corynebacteriaceae; Corynebacterium.
OX NCBI_TaxID=1718;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA Nakagawa S.;
RT "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF005277; BAB98444.1; -.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0008271; F:sulfate porter activity; IEA.
DR GO; GO:0008272; P:sulfate transport; IEA.

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DR InterPro; IPR006162; Ppantne S.
DR InterPro; IPR002645; STAS.
DR InterPro; IPR001902; Sulph_transpt.
DR Pfam; PF01740; STAS; 1.
DR Pfam; PF00916; Sulfate_transp; 1.
DR TIGRFAMs; TIGR00815; sulp; 1.
DR PROSITE; PS00012; PHOSPHOPANTHETHEINE; 1.
DR PROSITE; PS0801; STAS; 1.
KW Complete proteome.
SQ SEQUENCE 579 AA; 61660 MW; B70EA8BBE0C4C100 CRC64;

Query Match 57.3%; Score 43; DB 16; Length 579;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVGL 9
Db 48 AGLPAVVGL 56

RESULT 10
Q8FQM6 PRELIMINARY; PRT; 582 AA.
AC Q8FQM6;
DT 01-MAR-2003 (TREMBlrel. 23, Created)
DT 01-MAR-2003 (TREMBlrel. 23, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Putative sulfate transport protein.
GN CE1093.
OS Corynebacterium efficiens.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Corynebacteriaceae; Corynebacterium.
OX NCBI_TaxID=152794;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=YS-314 / AJ 12310 / DSM 44549 / JCM 11189;
RA Kawarabayasi Y., Yamazaki J., Hino Y., Kikuchi H., Nakamura Y.,
RA Ikeo K., Suzuki M., Mashima J., Itoh T., Yamagishi A., Nishio Y.,
RA Usuda Y., Sugimoto S.;
RT "The entire genomic sequence of Corynebacterium efficiens YS-314.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP005217; BAC17903.1; -.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0008271; F:sulfate porter activity; IEA.
DR GO; GO:0008272; P:sulfate transport; IEA.
DR InterPro; IPR002645; STAS.
DR InterPro; IPR001902; Sulph_transpt.
DR Pfam; PF01740; STAS; 1.
DR Pfam; PF00916; Sulfate_transp; 1.
DR TIGRFAMs; TIGR00815; sulp; 1.
DR PROSITE; PS0801; STAS; 1.
KW Complete proteome.
SQ SEQUENCE 582 AA; 62545 MW; CA67CFCABE04AE58 CRC64;

Query Match 57.3%; Score 43; DB 16; Length 582;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVGL 9
Db 51 AGLPAVVGL 59

RESULT 11
Q9HPR8 PRELIMINARY; PRT; 821 AA.
AC Q9HPR8;
DT 01-MAR-2001 (TREMBlrel. 16, Created)
DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE DNA helicase.
GN HEL OR VNG1501G.
OS Halobacterium sp. (strain NRC-1 / ATCC 700922 / JCM 11081).

```

```

Db      41 AGVPAIVVCETAGEAPTLSPAEQE 64
|||||:|||||:|||||
RESULT 13
Q89BH1 PRELIMINARY; PRT; 254 AA.
ID Q89BH1
AC Q89BH1;
CT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Bll18182 protein.
GN Bll18182.
OS Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiimi T.,
RA Sasamoto S., Watanabe A., Idegawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimo S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;
RT "Complete genomic sequence of nitrogen-fixing symbiotic bacterium
RT Bradyrhizobium japonicum USDA110."
RL DNA Res. 9:189-197(2002).
EMBL; AF005964; BAC53447.1; -.
DR InterPro; IPR002559; Transposase_11.
DR Pfam; PF01609; Transposase_11; 1.
KW Complete proteome.
SQ SEQUENCE 254 AA; 29160 MW; 76C1ED8F28D5C758 CRC64;

Query Match 56.0%; Score 42; DB 16; Length 254;
Best Local Similarity 66.7%; Pred. No. 98;
Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0

QY 2 GLPVGVLSPQE 13
|||||:|||||
DB 131 GLPVQLALSPQE 142

RESULT 14
Q9UXP0 PRELIMINARY; PRT; 326 AA.
ID Q9UXP0
AC Q9UXP0;
CT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE F420-dependent N5,N10-methylene-tetrahydromethanopterin reductase,
DE putative.
GN PFDA.
OS Methanolobus tindarius.
OC Archaea; Euryarchaeota; Euryarchaeota orders incertae sedis;
OC Methanosarcinales; Methanosarcinaceae; Methanolobus.
OX NCBI_TaxID=2221;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM 2278;
RX Westernberg D.J., Braune A., Ruppert C., Mueller V., Herzberg C.,
RA Gottschalk G., Blaut M.;
RT "The F420H2-dehydrogenase from Methanolobus tindarius: Cloning of the
RT ffd operon and expression of the genes in Escherichia coli.";
RT Submitted (SEP-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ011519; CAB56639.1; -.
DR FIR; T45226; T45226.
DR InterPro; IPR002103; Bac luciferase.
DR Pfam; PF00296; Bac luciferase; 1.
SQ SEQUENCE 326 AA; 34043 MW; 16F3AB9733A45D82 CRC64;

Query Match 56.0%; Score 42; DB 1; Length 326;
Best Local Similarity 57.1%; Pred. No. 1.3e+02;

```



Matches 8; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGLPAVVGGLSPGEG 14  
: |||||  
Db 84 SGGRAILGLPGEG 97

## RESULT 15

Q54756  
ID Q54756 PRELIMINARY; PRT; 574 AA.  
AC Q54756;  
DT 01-NOV-1996 (TREMELrel. 01, Created)  
DT 01-JUN-1998 (TREMELrel. 06, Last sequence update)  
DT 01-OCT-2003 (TREMELrel. 25, Last annotation update)  
DE Similar to plant sulfate transporter.  
OS Synechococcus sp. (strain PCC 7942) (Anacystis nidulans R2).  
OC Bacteria; Cyanobacteria; Chroococcales; Synechococcus.  
OX NCBI\_TaxID=1140;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=PCC 7942;  
RA Phung L.T., Haselkorn R.;  
RT "Gene encoding biotin carboxylase subunit of acetyl-CoA carboxylase  
from cyanobacterium Synechococcus sp. PCC 7942.";  
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.  
DR EMBL; U59234; AAB88215.1; -  
DR GO; GO:0016020; C:membrane; IEA.  
DR GO; GO:0005524; F:ATP binding; IEA.  
DR GO; GO:0004009; F:ATP-binding cassette (ABC) transporter acti...; IEA.  
DR GO; GO:0008271; F:sulfate porter activity; IEA.  
DR GO; GO:0008272; F:sulfate transport; IEA.  
DR GO; GO:0006810; F:transport; IEA.  
DR InterPro; IPR003439; ABC transporter.  
DR InterPro; IPR001902; Sulph\_transpt.  
DR Pfam; PF01740; STAS; 1.  
DR Pfam; PF00916; Sulfate\_transp; 1.  
DR TIGRFAMs; TIGR00815; sulp; 1.  
DR PROSITE; PS0801; STAS; 1.  
SQ SEQUENCE 574 AA; 62328 MW; 7BDECD65C67BBB00 CRC64;

Query Match 56.0%; Score 42; DB 2; Length 574;  
Best Local Similarity 88.9%; Pred. No. 2.3e+02;  
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVGGL 9  
: |||||  
Db 50 AGLPAVVGGL 58

Search completed: May 7, 2004, 12:37:57  
Job time : 29.2167 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: May 7, 2004, 12:19:40 ; Search time 41.85 seconds  
(without alignments)  
101.272 Million cell updates/sec

Title: US-09-786-214A-15

Perfect score: 75

Sequence: 1 AGLPAVWGLSPGEQE 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A\_Geneseq\_29Jan04.\*

1: Geneseqp1980s.\*

2: Geneseqp1990s.\*

3: Geneseqp2000s.\*

4: Geneseqp2001s.\*

5: Geneseqp2002s.\*

6: Geneseqp2003as.\*

7: Geneseqp2003bs.\*

8: Geneseqp2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	75	100.0	15	3	AY84269 Peptide d
2	75	100.0	20	3	AY84265 Truncated
3	75	100.0	25	3	AY84264 Peptide o
4	65	86.7	13	3	AY84268 Peptide d
5	65	86.7	14	3	AY84266 Peptide d
6	61	81.3	13	3	AY84267 Peptide d
7	45	60.0	234	4	AA836208 Human imm
8	44	58.7	116	4	ABG24038 Novel hum
9	43	57.3	473	4	AAU40252 Propionib
10	43	57.3	473	6	ABM36771 Propionib
11	43	57.3	565	4	AB76817 Coryneb
12	43	57.3	579	4	AA890917 C glutami
13	42	56.0	470	6	AAE34724 Streptomy
14	42	56.0	475	6	AAE34732 Streptomy
15	42	56.0	475	6	AAE34729 Streptomy
16	41	54.7	106	5	ABJ10397 Mutant an
17	41	54.7	106	5	ABJ10395 Mutant an
18	41	54.7	143	5	AAO21677 Human sec
19	41	54.7	143	7	ADB64633 Human pro
20	41	54.7	154	4	AAAM25921 Human pro
21	41	54.7	241	5	AAAM48925 scfv anti
22	41	54.7	294	1	AAAP82484 Tropoelas
23	41	54.7	370	2	AAAY14924 Amino aci
24	41	54.7	370	6	ABP70895 Mycobacte
25	41	54.7	372	5	ABB73530 M vaccae

26	41	54.7	396	3	AB51784 Gene 15 h
27	41	54.7	530	7	ADD49105 Human NOV
28	41	54.7	549	7	ADD49091 Human NOV
29	41	54.7	579	5	ABG70277 Human Gly
30	41	54.7	579	5	ABG97356 Human CGD
31	41	54.7	579	6	ABR39111 Human GPC
32	41	54.7	579	7	ADD49087 Human NOV
33	41	54.7	579	7	ADD49107 Human NOV
34	41	54.7	579	7	ADD49089 Human NOV
35	41	54.7	592	7	ADD49099 Human NOV
36	41	54.7	774	3	ABO0077 Human lys
37	41	54.7	774	5	ABO07649 Human LOR
38	41	54.7	774	6	ABU03509 Angiogene
39	41	54.7	774	6	ABU03509 Angiogene
40	41	54.7	774	6	ABU03509 Angiogene
41	41	54.7	11096	4	AAE10129 Streptomy
42	40	53.3	119	4	AAU51147 Propionib
43	40	53.3	119	6	ABM47666 Propionib
44	40	53.3	132	3	AA836582 Arabidops
45	40	53.3	215	5	ABB97875 Human sec

## ALIGNMENTS

### RESULT 1

AY84269	12-JUL-2000 (first entry)	Peptide derived from macrophage colony stimulating gene alternative ORF.
ID	AY84269 standard; peptide; 15 AA.	tumour rejection antigen; macrophage colony stimulating gene;
XX	AY84269;	macrophage-colony stimulating factor; antigen presenting cell;
AC		human leukocyte antigen; CD8+ cytotoxic T lymphocyte.
XX		Synthetic.
DT		Homo sapiens.
XX		WO200013699-A1.
DE		16-MAR-2000.
XX		03-SEP-1999; 99WO-US020344.
KW		04-SEP-1998; 98US-0099077P.
XX		(LUDW-) LUDWIG INST CANCER RES.
XX		Probst-Keppler M, Van Den Eynde B, Boon-Falleur T;
XX		WPI; 2000-256859/22.
XX		Isolated polypeptide used to treat subjects having a disorder
XX		characterized by expression of alternative open reading frame macrophage-
XX		colony stimulating factor comprises 25 amino acid residue sequence.
XX		Example 2; Page 40; 74pp; English.
XX		The present sequence represents a peptide which is derived from a tumour
XX		rejection antigen precursor encoded by an alternative open reading frame
XX		(ORF) of human macrophage colony stimulating gene. Peptides derived from
XX		the alternative ORF of macrophage-colony stimulating factor, when
XX		presented by an antigen presenting cell having a human leukocyte antigen
XX		(HLA) class I molecule, effectively induce the activation and
XX		proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic
XX		acids derived from the alternate ORF of macrophage-colony stimulating
XX		factor are useful for enriching selectively a population of T lymphocytes
XX		with CD8+ T lymphocytes. They are also useful for diagnosing a disorder
XX		characterized by expression of the polypeptide, and for identifying

CC functional variants and mimetics

SQ Sequence 15 AA;

Query Match 100.0%; Score 75; DB 3; Length 15;  
Best Local Similarity 100.0%; Pred. No. 4.7e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVLSPGGEQ 15  
| | | | | | | | | | | | | | |  
Db 1 AGLPAVVLSPGGEQ 15

RESULT 2

AY84265  
ID AAY84265 standard; peptide; 20 AA.

XX AC AAY84265;

XX 12-JUL-2000 (first entry)

XX Truncated macrophage colony stimulating factor tumour antigen.

XX tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX OS Homo sapiens.

XX PN WO200013699-A1.

XX PD 16-MAR-2000.

XX PF 03-SEP-1999; 99WO-US020344.

XX PR 04-SEP-1998; 98US-0099077P.

XX PA (LUDW-) LUDWIG INST CANCER RES.

XX PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX DR N-PSDB; AAZ99675.

XX Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 3; Page 64; 74pp; English.

XX The present sequence represents a truncated tumour rejection antigen  
CC precursor, and is encoded by a truncated alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX SQ Sequence 20 AA;

Query Match 100.0%; Score 75; DB 3; Length 20;  
Best Local Similarity 100.0%; Pred. No. 6.5e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVLSPGGEQ 15  
| | | | | | | | | | | | | | |  
Db 2 AGLPAVVLSPGGEQ 16

RESULT 3

AY84264  
ID AAY84264 standard; peptide; 25 AA.

XX AC AAY84264;

XX 12-JUL-2000 (first entry)

XX Peptide of alternate reading frame of macrophage colony stimulating gene.  
KW Renal cell carcinoma; antigen; cytotoxic T lymphocyte;  
KW tumour rejection antigen; macrophage colony stimulating gene;  
KW macrophage-colony stimulating factor; antigen presenting cell;  
KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX OS Homo sapiens.

XX PN WO200013699-A1.

XX PD 16-MAR-2000.

XX PF 03-SEP-1999; 99WO-US020344.

XX PR 04-SEP-1998; 98US-0099077P.

XX PA (LUDW-) LUDWIG INST CANCER RES.

XX PI Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX DR N-PSDB; AAZ99672.

XX Isolated polypeptide used to treat subjects having a disorder  
PT characterized by expression of alternative open reading frame macrophage-  
PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 1; Page 64; 74pp; English.

XX The present sequence represents a tumour rejection antigen precursor, and  
CC is encoded by an alternative open reading frame (ORF) of human macrophage  
CC colony stimulating gene. Peptides derived from the alternative ORF of  
CC macrophage-colony stimulating factor, when presented by an antigen  
CC presenting cell having a human leukocyte antigen (HLA) class I molecule,  
CC effectively induce the activation and proliferation of CD8+ cytotoxic T  
CC lymphocytes. Polypeptide and nucleic acids derived from the alternate ORF  
CC of macrophage-colony stimulating factor are useful for enriching  
CC selectively a population of T lymphocytes with CD8+ T lymphocytes. They  
CC are also useful for diagnosing a disorder characterized by expression of  
CC the polypeptide, and for identifying functional variants and mimetics

XX SQ Sequence 25 AA;

Query Match 100.0%; Score 75; DB 3; Length 25;  
Best Local Similarity 100.0%; Pred. No. 8.2e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVLSPGGEQ 15  
| | | | | | | | | | | | | | |  
Db 2 AGLPAVVLSPGGEQ 16

RESULT 4

AY84268

ID AAY84268 standard; peptide; 13 AA.

XX AC AAY84268;

XX 12-JUL-2000 (first entry)

XX Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;

KW macrophage-colony stimulating factor; antigen presenting cell;  
 XX human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Synthetic.  
 OS Homo sapiens.

FN WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 86.7%; Score 65; DB 3; Length 13;  
 Best Local Similarity 100.0%; Pred. No. 0.0015;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LPAVGLSPGEQE 15

Db 1 LPAVGLSPGEQE 13

RESULT 5

AAAY84266  
 ID AAY84266 standard; peptide; 14 AA.

AC AAY84266;

DT 12-JUL-2000 (first entry)

DE Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

XX Synthetic.

OS Homo sapiens.

FN WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX

PR 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

XX Claim 2; Page 39; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
 CC rejection antigen precursor encoded by an alternative open reading frame  
 CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
 CC the alternative ORF of macrophage-colony stimulating factor, when  
 CC presented by an antigen presenting cell having a human leukocyte antigen  
 CC (HLA) class I molecule, effectively induce the activation and  
 CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
 CC acids derived from the alternate ORF of macrophage-colony stimulating  
 CC factor are useful for enriching selectively a population of T lymphocytes  
 CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
 CC characterized by expression of the polypeptide, and for identifying  
 CC functional variants and mimetics

XX Sequence 14 AA;

Query Match 86.7%; Score 65; DB 3; Length 14;  
 Best Local Similarity 100.0%; Pred. No. 0.0017;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LPAVGLSPGEQE 15

Db 1 LPAVGLSPGEQE 13

RESULT 6

AAAY84267  
 ID AAY84267 standard; peptide; 13 AA.

AC AAY84267;

DT 12-JUL-2000 (first entry)

DE Peptide derived from macrophage colony stimulating gene alternative ORF.

XX tumour rejection antigen; macrophage colony stimulating gene;  
 KW macrophage-colony stimulating factor; antigen presenting cell;  
 KW human leukocyte antigen; CD8+ cytotoxic T lymphocyte.

OS Synthetic.

OS Homo sapiens.

FN WO200013699-A1.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-US020344.

XX 04-SEP-1998; 98US-0099077P.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Probst-Kepper M, Van Den Eynde B, Boon-Falleur T;

XX WPI; 2000-256859/22.

XX Isolated polypeptide used to treat subjects having a disorder  
 PT characterized by expression of alternative open reading frame macrophage-  
 PT colony stimulating factor comprises 25 amino acid residue sequence.

PS Example 2; Page 40; 74pp; English.

XX The present sequence represents a peptide which is derived from a tumour  
CC rejection antigen precursor encoded by an alternative open reading frame  
CC (ORF) of human macrophage colony stimulating gene. Peptides derived from  
CC the alternative ORF of macrophage-colony stimulating factor, when  
CC presented by an antigen presenting cell having a human leukocyte antigen  
CC (HLA) class I molecule, effectively induce the activation and  
CC proliferation of CD8+ cytotoxic T lymphocytes. Polypeptide and nucleic  
CC acids derived from the alternate ORF of macrophage-colony stimulating  
CC factor are useful for enriching selectively a population of T lymphocytes  
CC with CD8+ T lymphocytes. They are also useful for diagnosing a disorder  
CC characterized by expression of the polypeptide, and for identifying  
CC functional variants and mimetics

XX Sequence 13 AA;

Query Match 81.3%; Score 61; DB 3; Length 13;  
Best Local Similarity 100.0%; Pred. No. 0.0066;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 PAVVGLSPGEQ 15  
||:|||||  
Db 1 PAVVGLSPGEQ 12

RESULT 7

AAB36208  
ID AAB36208 standard; protein; 234 AA.

AC AAB36208;

DT 15-FEB-2001 (first entry)

DE Human immune system associated protein HISAP-6.

XX Human; immune system associated protein; HISAP-6; immune disorder;  
KW infection; autoimmune disease; cancer.

XX Homo sapiens.

PN US6135941-A.

XX 24-OCT-2000.

XX 27-MAR-1998; 98US-00049672.

XX 27-MAR-1998; 98US-00049672.

XX (INCY-) INCYTE PHARM INC.

XX Tang YT, Yue H, Lal P, Corley NC, Guegler KJ, Baughn MR;  
PI Hillman JL, Au-Young J;

XX WPI: 2001-030926/04.

XX N-PSDB; AAC66524.

XX New human immune system associated proteins (HISAP) and polynucleotides  
PT encoding the HISAP, useful for diagnosing, treating or preventing immune  
PT or cell proliferative disorders or infections.

XX Claim 1; Col 59-60; 54pp; English.

XX The present invention provides the coding and protein sequences for a  
CC number of human immune system associated proteins (HISAPs). These can be  
CC used in the diagnosis and treatment of various autoimmune disorders,  
CC infections and cell proliferation diseases. The diseases include AIDS,  
CC adult respiratory distress syndrome, anaemia, asthma, atherosclerosis,  
CC Crohn's disease, irritable bowel syndrome, multiple sclerosis, myasthenia  
CC gravis, osteoarthritis, rheumatoid arthritis, scleroderma, systemic lupus  
CC erythematosus, arteriosclerosis, cirrhosis and cancer

XX Sequence 234 AA;

Query Match

Best Local Similarity 60.0%; Score 45; DB 4; Length 234;  
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 4 PAVVGLSPGEQ 14  
||:|||||  
Db 28 PAVVGLSPGER 38

RESULT 8

ABG24038  
ID ABG24038 standard; protein; 116 AA.

XX ABG24038;

DT 18-FEB-2002 (first entry)

DE Novel human diagnostic protein #24029.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder.

XX Homo sapiens.

PN WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US008631.

XX 31-MAR-2000; 2000US-00540217.

XX 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI: 2001-639362/73.

XX N-PSDB; AAS88225.

XX New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity.

XX Claim 20; SEQ ID NO 54397; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypeptide (II)  
CC sequences. (I) is useful as hybridisation probes, polymerase chain  
CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,  
CC and in recombinant production of (II). The polynucleotides are also used  
CC in diagnostics as expressed sequence tags for identifying expressed  
CC genes. (I) is useful in gene therapy techniques to restore normal  
CC activity of (II) or to treat disease states involving (II). (II) is  
CC useful for generating antibodies against it, detecting or quantitating a  
CC polypeptide in tissue, as molecular weight markers and as a food  
CC supplement. (II) and its binding partners are useful in medical imaging  
CC of sites expressing (II). (I) and (II) are useful for treating disorders  
CC involving aberrant protein expression or biological activity. The  
CC polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. ABG00010-ABG03077 represent novel human diagnostic  
CC amino acid sequences of the invention. Note: The sequence data for this  
CC patent did not appear in the printed specification, but was obtained in  
CC electronic format directly from WIPO at  
CC ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 116 AA;

Query Match

58.7%; Score 44; DB 4; Length 116;

Best Local Similarity 64.3%; Pred. No. 35;  
Matches 9; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 GLPAVVGSLSPGEOR 15  
|||:|||||:|  
Db 90 GLPXURGLSPVERE 103

RESULT 9  
AAU40252  
ID AAU40252 standard; protein; 473 AA.  
XX  
AC AAU40252;  
XX  
DT 13-FEB-2002 (first entry)  
XX  
DE Propionibacterium acnes immunogenic protein #1148.  
XX  
KW SAPHO syndrome, synovitis; acne; pustulosis; hypertosis; osteomyelitis;  
XX uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;  
XX inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
XX dermatological; osteopathic; neuroprotectant.  
XX  
OS Propionibacterium acnes.  
XX  
FN WO200181581-A2.  
XX  
PD 01-NOV-2001.  
XX  
PF 20-APR-2001; 2001WO-US012865.  
XX  
PR 21-APR-2000; 2000US-0199047P.  
XX  
PR 02-JUN-2000; 2000US-0208841P.  
XX  
PR 07-JUL-2000; 2000US-0216747P.  
XX  
PA (CORI-) CORIXA CORP.  
XX  
PI Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;  
PI L'maisonneuve J, Zhang Y, Jen S, Carter D;  
XX  
XX  
DR WPI: 2001-616774/71.  
DR N-PSDB; AAS59511.  
XX  
FT Propionibacterium acnes polypeptides and nucleic acids useful for  
FT vaccinating against and diagnosing infections, especially useful for  
FT treating acne vulgaris.  
XX  
PS Example 1; SEQ ID NO 1447; 1069pp; English.  
XX  
CC Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic  
CC polypeptides. The proteins and their associated DNA sequences are used in  
CC the treatment, prevention and diagnosis of medical conditions caused by  
CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,  
CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.  
CC P. acnes is also involved in infections of bone, joints and the central  
CC nervous system, however it is particularly involved in the inflammatory  
CC lesions associated with acne vulgaris. A method for detecting the  
CC presence or absence of P. acnes in a patient comprises contacting a  
CC sample with a binding agent that binds to the proteins of the invention  
CC and determining the amount of bound protein in the sample. The  
CC polypeptides may be used as antigens in the production of antibodies  
CC specific for P. acnes proteins. These antibodies can be used to  
CC downregulate expression and activity of P. acnes polypeptides and  
CC therefore treat P. acnes infections. The antibodies may also be used as  
CC diagnostic agents for determining P. acnes presence, for example, by  
CC enzyme linked immunosorbent assay (ELISA). Note: The sequence data for  
CC this patent did not form part of the printed specification, but was  
CC obtained in electronic format directly from WIPO at  
CC ftp.wipo.int/pub/published\_pct\_sequences  
XX  
SQ Sequence 473 AA;

Best Local Similarity 63.6%; Pred. No. 2.3e+02;  
Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GLPAVVGSLSPG 12  
|||:|||||:|  
Db 446 GLPAIIGLAAG 456

RESULT 10  
ABM36771  
ID ABM36771 standard; protein; 473 AA.  
XX  
AC ABM36771;  
XX  
DT 20-OCT-2003 (first entry)  
XX  
DE Propionibacterium acnes predicted ORF-encoded polypeptide #1447.  
XX  
KW Acne vulgaris; antiseborrheic; dermatological; antibacterial;  
XX immunostimulant; immune response; vaccine.  
XX  
OS Propionibacterium acnes.  
XX  
FN WO2003033515-A1.  
XX  
PD 24-APR-2003.  
XX  
PF 11-OCT-2002; 2002WO-US032727.  
XX  
PR 15-OCT-2001; 2001US-00978925.  
XX  
PA (CORI-) CORIXA CORP.  
XX  
PI Mitcham JL, Skeiky YAW, Persing DH, Bhatia A, Maisonneuve JL;  
PI Zhang Y, Wang S, Jen S, Lodes MJ, Benson DR, Jones R, Carter D;  
PI Barth B, Vallie-Douglass J;  
XX  
XX WPI: 2003-381789/36.  
DR N-PSDB; ACF64440.  
XX  
FT New Propionibacterium acnes polypeptides and polynucleotides encoding the  
FT polypeptide, useful for diagnosing, preventing or treating acne vulgaris,  
FT or for stimulating an immune response specific for a P. acnes protein.  
XX  
PS Example 1; SEQ ID NO 1447; 1481pp; English.  
XX  
CC The invention relates to an isolated polynucleotide (ACF64435-ACF64733)  
CC encoding a Propionibacterium acnes protein. The invention also relates to  
CC polypeptides encoded by the polynucleotides (ABM35624-ABM64536) and to  
CC immunogenic fragments of P. acnes polypeptides. The invention  
CC additionally encompasses expression vectors and host cells comprising a  
CC polynucleotide of the invention; antibodies against polypeptides of the  
CC invention; fusion proteins comprising a polypeptide of the invention; a  
CC method for stimulating an immune response specific for a P. acnes  
CC polypeptide and an isolated T cell population comprising T cells prepared  
CC via this method; a vaccine composition (comprising P. acnes polypeptides,  
CC polynucleotides, antibodies, fusion proteins, T cell populations, or  
CC antigen-presenting cells that express the polypeptide); a method and kit  
CC for detecting or determining the presence or absence of P. acnes in a  
CC patient; and a method for inhibiting the development of P. acnes in a  
CC patient. The P. acnes polypeptides, polynucleotides, antibodies, fusion  
CC proteins, T cell populations or antigen-presenting cells that express the  
CC polypeptides are useful for diagnosing, preventing or treating acne  
CC vulgaris, or for stimulating an immune response specific for a P. acnes  
CC protein. The polynucleotides can also be used as probes or primers for  
CC nucleic acid hybridisation. The vaccine composition is useful for the  
CC stimulation of an immune response against P. acnes, or for treating acne,  
CC and the kit is useful for performing a diagnostic assay. The present  
CC sequence represents a polypeptide predicted to be encoded by an ORF (open  
CC reading frame) contained within the P. acnes polynucleotides of the  
CC invention. Note: The sequence data for this patent did not form part of  
CC the printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
CC

XX SQ Sequence 473 AA;  
 Query Match 57.3%; Score 43; DB 6; Length 473;  
 Best Local Similarity 63.6%; Pred. No. 2.3e+02;  
 Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;  
 QY 2 GLPAVGLSPG 12  
 Db 446 GLPAIIGLAAG 456  
 RESULT 11  
 AAB76817  
 ID AAB76817 standard; protein; 565 AA.  
 AC AAB76817;  
 DT 11-APR-2001 (first entry)  
 DE Corynebacterium glutamicum MCT protein SEQ ID NO:616.  
 XX  
 KW Corynebacterium glutamicum; brevibacterium lactofermentum; MCT;  
 KW membrane construction and membrane transport protein; petroleum spill;  
 KW hydrocarbon degradation; gram positive aerobic bacterium; marker;  
 KW identification; microorganism; fine chemical production; transformation;  
 KW genome mapping; genetic engineering.  
 XX  
 OS Corynebacterium glutamicum.  
 XX  
 PN WO200100805-A2.  
 XX  
 PD 04-JAN-2001.  
 XX  
 PF 23-JUN-2000; 2000WO-IB000926.  
 XX  
 PR 25-JUN-1999; 99US-0141031P.  
 PR 08-JUL-1999; 99DE-0103145A.  
 PR 08-JUL-1999; 99DE-0103147B.  
 PR 08-JUL-1999; 99DE-01031563.  
 PR 09-JUL-1999; 99DE-01032122.  
 PR 09-JUL-1999; 99DE-01032124.  
 PR 09-JUL-1999; 99DE-01032125.  
 PR 09-JUL-1999; 99DE-01032128.  
 PR 09-JUL-1999; 99DE-01032180.  
 PR 09-JUL-1999; 99DE-01032182.  
 PR 09-JUL-1999; 99DE-01032190.  
 PR 09-JUL-1999; 99DE-01032191.  
 PR 09-JUL-1999; 99DE-01032205.  
 PR 09-JUL-1999; 99DE-01032212.  
 PR 09-JUL-1999; 99DE-01032227.  
 PR 09-JUL-1999; 99DE-01032228.  
 PR 09-JUL-1999; 99DE-01032229.  
 PR 09-JUL-1999; 99DE-01032230.  
 PR 14-JUL-1999; 99DE-01032927.  
 PR 14-JUL-1999; 99DE-01033005.  
 PR 14-JUL-1999; 99DE-01033006.  
 PR 27-AUG-1999; 99DE-01040764.  
 PR 27-AUG-1999; 99DE-01040765.  
 PR 27-AUG-1999; 99DE-01040766.  
 PR 27-AUG-1999; 99DE-01040830.  
 PR 27-AUG-1999; 99DE-01040831.  
 PR 27-AUG-1999; 99DE-01040832.  
 PR 27-AUG-1999; 99DE-01040833.  
 PR 31-AUG-1999; 99DE-01041378.  
 PR 31-AUG-1999; 99DE-01041379.  
 PR 31-AUG-1999; 99DE-01041395.  
 PR 03-SEP-1999; 99DE-01042077.  
 PR 03-SEP-1999; 99DE-01042078.  
 PR 03-SEP-1999; 99DE-01042079.  
 PR 03-SEP-1999; 99DE-01042088.  
 XX  
 (BADI ) BASF AG.

XX PI Pompejus M, Kroeger B, Schroeder H, Zelder O, Haberhauer G;  
 XX  
 DR WPI; 2001-071486/08.  
 DR N-PSDB; AAF68050.  
 XX  
 PT Corynebacterium glutamicum nucleic acids encoding membrane construction  
 PT and membrane transport proteins or their portions, useful for typing or  
 PT identifying C. glutamicum or related bacteria, and as markers for  
 PT transformation.  
 XX  
 PS Claim 20; Page 1030-1031; 1119pp; English.  
 XX  
 CC AAF67743 to AAF68080 encode the Corynebacterium glutamicum membrane  
 CC construction and membrane transport (MCT) proteins given in AAB76510 to  
 CC AAB76847. The MCT nucleic acids and proteins are useful in the  
 CC identification of microorganisms which can be used to produce fine  
 CC chemicals, for modulating fine chemical production in C. glutamicum or  
 CC related bacteria (e.g. Brevibacterium lactofermentum), the typing or  
 CC identification of C. glutamicum or related bacteria, as reference points  
 CC for mapping C. glutamicum genome, and as markers for transformation.  
 CC AAF68082 and AAF68082 represent sequencing primers which are used in an  
 CC example from the present invention  
 XX  
 SQ Sequence 565 AA;  
 Query Match 57.3%; Score 43; DB 4; Length 565;  
 Best Local Similarity 100.0%; Pred. No. 2.8e+02;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 AGLPAVGL 9  
 Db 34 AGLPAVGL 42  
 RESULT 12  
 AAG90917  
 ID AAG90917 standard; protein; 579 AA.  
 XX  
 AC AAG90917;  
 XX  
 DT 26-SEP-2001 (first entry)  
 DE C glutamicum protein fragment SEQ ID NO: 4671.  
 XX  
 KW Corynebacterium; amino acid synthesis; vitamin; saccharide;  
 KW organic acid synthesis.  
 XX  
 OS Corynebacterium glutamicum.  
 XX  
 PN EP1108790-A2.  
 XX  
 PD 20-JUN-2001.  
 XX  
 PF 18-DEC-2000; 2000EP-00127688.  
 PR 16-DEC-1999; 99JP-00377484.  
 PR 07-APR-2000; 2000JP-00159162.  
 PR 03-AUG-2000; 2000JP-00280988.  
 XX  
 PA (KYOW ) KYOWA HAKKO KOGYO KK.  
 XX  
 PI Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;  
 PI Tateishi N, Senoh A, Ikeda M, Ozaki A;  
 XX  
 DR WPI; 2001-376931/40.  
 DR N-PSDB; AAF66136.  
 XX  
 PT Novel polynucleotides derived from Corynebacterium bacteria, for identifying  
 PT mutation point of a gene, measuring expression of a gene, analyzing  
 PT expression profile or pattern of a gene and identifying homologous gene.  
 XX  
 PS Claim 17; SEQ ID NO 4671; 246pp + Sequence Listing; English.

XX The present invention provides a number of nucleotide and protein  
 CC sequences from the Coryneform bacterium *Corynebacterium glutamicum*. These  
 CC are useful for identifying the mutation point of a gene derived from a  
 CC mutant of coryneform bacterium, measuring expression amount and analysing  
 CC the expression profile or expression pattern of a gene derived from  
 CC coryneform bacterium, and identifying a homologue of a gene derived from  
 CC coryneform bacterium. Coryneform bacteria are useful for producing amino  
 CC acids, nucleic acids, vitamins, saccharides and organic acids,  
 CC particularly L-lysine. The present sequence is a protein described in the  
 CC exemplification of the invention. Note: The sequence data for this patent  
 CC did not form part of the printed specification, but was obtained in  
 CC electronic format directly from the European Patent Office

XX SQ Sequence 579 AA;

Query Match 57.3%; Score 43; DB 4; Length 579;  
 Best Local Similarity 100.0%; Pred. No. 2.9e+02;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGLPAVVG 9  
 Db 48 AGLPAVVG 56

RESULT 13  
 AAE34724  
 ID AAE34724 standard; protein; 470 AA.

XX AC AAE34724;

XX DT 14-MAY-2003 (first entry)

XX DE Streptomyces rimosus ema3 protein.

XX KW P450 monooxygenase; avermectin; ferrodioxin; ferrodioxin reductase; enzyme;  
 KW emamectin; insecticide; ema3 protein.

XX OS Streptomyces rimosus.

XX PN WO200292801-A2.

XX PD 21-NOV-2002.

XX PF 15-MAY-2002; 2002WO-EP005363.

XX PR 16-MAY-2001; 2001US-0291149P.

XX PA (SYGN ) SYNGENTA PARTICIPATIONS AG.

XX PI Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;

XX DR WPI; 2003-140280/13.  
 DR N-PSDB; AAD53019.

XX PT Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.

XX PS Claim 17; Page 107-108; 157pp; English.

XX The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodioxins and ferrodioxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus ema3 protein

XX SQ Sequence 470 AA;

Query Match 56.0%; Score 42; DB 6; Length 470;  
 Best Local Similarity 72.7%; Pred. No. 3.3e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 3 LPVVVGLSPGE 13  
 Db 44 LPSVVGHPGE 54

RESULT 14  
 AAE34732

ID AAE34732 standard; protein; 475 AA.

XX AC AAE34732;

XX DT 14-MAY-2003 (first entry)

XX DE Streptomyces rimosus ema1 protein.

XX KW P450 monooxygenase; avermectin; ferrodioxin; ferrodioxin reductase; enzyme;  
 KW emamectin; insecticide; ema1 protein.

XX OS Streptomyces rimosus.

XX PN WO200292801-A2.

XX PD 21-NOV-2002.

XX PF 15-MAY-2002; 2002WO-EP005363.

XX PR 16-MAY-2001; 2001US-0291149P.

XX PA (SYGN ) SYNGENTA PARTICIPATIONS AG.

XX PI Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
 PI Buckel TG;

XX DR WPI; 2003-140280/13.  
 DR N-PSDB; AAD53027.

XX PT Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
 PT and capable of regioselectively oxidizing alcohol of avermectin useful  
 PT for making emamectin from avermectin.

XX PS Claim 17; Page 121-122; 157pp; English.

XX The present invention relates to novel proteins that exhibit an enzymatic  
 CC activity of P450 monooxygenase and capable of regioselectively oxidising  
 CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
 CC The invention also relates to ferrodioxins and ferrodioxin reductases that  
 CC are active with the P450 monooxygenases. Sequences of the invention are  
 CC useful for producing 4-keto-avermectin from avermectin, which is useful  
 CC for producing emamectin. Emamectin is useful as an insecticide. The  
 CC present sequence is Streptomyces rimosus ema1 protein

XX SQ Sequence 475 AA;

Query Match 56.0%; Score 42; DB 6; Length 475;  
 Best Local Similarity 72.7%; Pred. No. 3.3e+02;  
 Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 3 LPVVVGLSPGE 13  
 Db 49 LPSVVGHPGE 59

RESULT 15  
 AAE34729

ID AAE34729 standard; protein; 475 AA.

XX AC AAE34729;

XX DT 14-MAY-2003 (first entry)



XX Streptomyces albofaciens ema8 protein.  
DE  
XX P450 monooxygenase; avermectin; ferrodoxin; ferrodoxin reductase; enzyme;  
KW emamectin; insecticide; ema8 protein.  
KW  
XX Streptomyces albofaciens.  
OS  
XX WO200292801-A2.  
PN  
XX  
XX 21-NOV-2002.  
PD  
XX  
XX 15-MAY-2002; 2002WO-BF005363.  
PF  
XX  
XX 16-MAY-2001; 2001US-0291149P.  
PR  
XX (SYGN ) SYNGENTA PARTICIPATIONS AG.  
PA  
XX Molnar I, Ligon JM, Zirkle RE, Hammer PE, Hill DS, Pachlatko JP;  
PI Buckel TG;  
PI  
XX  
XX WPI; 2003-140280/13.  
DR  
XX N-PSDB; AAD53024.  
DR  
XX Novel polypeptide that exhibits enzymatic activity of P450 monooxygenase  
PT and capable of regioselectively oxidizing alcohol of avermectin useful  
PT for making emamectin from avermectin.  
PT  
XX  
XX Claim 17; Page 116-117; 157pp; English.  
PS  
XX The present invention relates to novel proteins that exhibit an enzymatic  
CC activity of P450 monooxygenase and capable of regioselectively oxidising  
CC the 4-carbinol group at position 4 of avermectin to 4-keto-avermectin.  
CC The invention also relates to ferrodoxins and ferrodoxin reductases that  
CC are active with the P450 monooxygenases. Sequences of the invention are  
CC useful for producing 4-keto-avermectin from avermectin, which is useful  
CC for producing emamectin. Emamectin is useful as an insecticide. The  
CC present sequence is Streptomyces albofaciens ema8 protein  
XX  
SQ Sequence 475 AA;

Query Match 56.0%; Score 42; DB 6; Length 475;  
Best Local Similarity 72.7%; Pred. No. 3.3e+02;  
Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
QY 3 LPAVVGLSPGE 13  
Db 49 LPSYVGLHPGE 59  
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||: ||| |||

Search completed: May 7, 2004, 12:33:48  
Job time : 43.85 secs